

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁵ : C07D 233/70, A61K 31/415 C07D 233/84, 235/02, 403/10 C07D 471/10, 491/10, 495/10	A1	(11) International Publication Number: WO 93/040 (43) International Publication Date: 4 March 1993 (04.03.)
(21) International Application Number: PCT/U (22) International Filing Date: 19 August 1992 (30) Priority data: 19 August 1991 (19.08.9 929,455 14 August 1992 (14.08.9 14.08.9 19.08) 19.08 (19.08.9 19.08) 19.09 (19.08.9 19.08) 19.09 (19.08.9 19.09) 19.08 (19.08.9 19.09) 19.09 (19.08.9	(19.08. 1) 2) ND COngton, 1 ord Rondawat 810 (U	mours and Company, Legal/Patent Records Cent 1007 Market Street, Wilmington, DE 19898 (US). [81] Designated States: AU, CA, CS, JP, KR, PL, European tent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, LU, MC, NL, SE). [8] [9] [9] [9] [10]

(54) Title: ANGIOTENSIN II RECEPTOR BLOCKING IMIDAZOLINONE DERIVATIVES

(57) Abstract

Novel imidazolinone derivatives of formula (I), which are useful as angiotensin II antagonists, are disclosed.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FI	Finland	MN	Mongolia
AU	Australia	FR	France	MR	Mauritania
BB	Barbados	GA	Gabon	MW	Malawi
86	Belgium	GB	United Kingdom	NL	Netherlands
8F	Burkina Faso	GN	Guinca	NO	Norway
BG	Bulgaria	GR	Greece	NZ	New Zealand
BJ	Benin	HU	Hungary	PL.	Poland
BR	Brazil	1E	Ireland	PT	Portugul
CA	Canada	IT	Haly	RO	Romania
CF	Central African Republic	JP	Lapan	RU	Russian Federatinn
		KP	Democratic People's Republic	SD	Sudan
CC	Congo	Kr	of Korea	SE	Sweden
CH	Switzerland	KR	Republic of Korea	SK	Slovak Republic
CI	Côte d'Ivoire	Li	Liechtenstein	SN	Senegal
CM	Cameroun			SU	Soviet Union
CS	Ozerhoslovakia	LK	Sri Lanka	TD	Chad
CZ	Czech Republic	LU	Luxembourg	TG	
DE	Germany	MC	Monacu		Togo
DK	Denmark	MG	Madagascar	UA	Ukraine
FS	South	MI	Mali	US	United States of America

WO 93/04046 PCT/US92/07022

1

TITLE

ANGIOTENSIN II RECEPTOR BLOCKING IMIDAZOLINONE DERIVATIVES

CROSS-REFERENCE TO RELATED APPLICATION

This application is a continuation-in-part of U.S. Application Serial Number 07/747,023, filed August 19, 10 1991.

BACKGROUND OF THE INVENTION

Field of the Invention

15

20

5

This invention relates to novel substituted imidazolinone derivatives. The invention also relates to pharmaceutical compositions containing the novel imidazolinone derivatives and pharmaceutical methods using them, alone and in conjugation with other drugs.

The compounds of this invention inhibit the action of the hormone angiotensin II (AII) and are useful therefore in alleviating angiotensin induced hypertension. The enzyme renin acts on a blood plasma a2-globulin, angiotensinogen, to produce angiotensin I, which is then converted by ACE to AII. The latter substance is a powerful vasopressor agent which has been implicated as a causative agent for producing high blood pressure in various mammalian species, such as the rat, dog, and man. The compounds of this invention inhibit the action of AII at its receptors on target cells and thus prevent the increase in blood pressure produced by this hormone-receptor interaction. By administering a compound of this invention to a species of mammal with

15

20

25

30

hypertension due to AII, the blood pressure is reduced. The compounds of this invention are also useful for the treatment of congestive heart failure. Administration of a compound of this invention with a diuretic such as furosemide or hydrochlorothiazide, either as a stepwise combined therapy (diuretic first) or as a physical mixture, enhances the antihypertensive effect of the compound. Administration of a compound of this invention with a NSAID can prevent renal failure which sometimes results from administration of a NSAID.

Several peptide analogs of AII are known to inhibit the effects of this hormone by competitively blocking the receptors, but their experimental and clinical applications have been limited by the partial agonist activity and lack of oral absorption (M. Antonaccio, Clin. Exp. Hypertens., 1982, A4, 27-46; D. H. P. Streeten and G. H. Anderson, Jr., Handbook of Hypertension. Clinical Pharmacology of Antihypertensive Drugs, ed., A. E. Doyle, Vol. 5, pages 246-271, Elsevier Science Publisher, Amsterdam, The Netherlands, 1984).

Several non-peptide antagonists of AII have been disclosed. These compounds are covered by U.S. Patents 4,207,324; 4,340,598; 4,576,958; 4,582,847; and 4,880,804; in European Patent Applications 028,834; 245,637; 253,310; and 291,969; and in articles by A. T. Chiu, et al. (Eur. J. Pharm. Exp. Therap., 1988, 157, 13-21) and by P. C. Wong, et al. (J. Pharm. Exp. Therap, 1988, 247, 1-7). All of the U.S. Patents, European Patent Applications 028,834 and 253,310 and the two articles disclose substituted imidazole compounds which are generally bonded through a lower alkyl bridge to a substituted phenyl. European Patent Application 245,637 discloses derivatives of 4,5,6,7-tetrahydro-2H-imidazo(4,5-g)pyridine-6-carboxylic acid and analogs

thereof as antihypertensive agents, specifically ${\tt Ca}^{2+}$ channel blockers.

L. Chang et al., in EP 0 412 594 A (filed July 23, 1990) disclose substituted triazolinones,

5 triazolinethiones, and triazolinimines of the formula:

These are claimed to be antagonists of AII which are 10 useful for treating hypertension, congestive heart failure (CHF), and elevated intraocular pressure.

C. Bernhart et al., in WO 91/14679 (published October 3, 1991) disclose heterocyclic N-substituted derivatives of the formula

WO 93/04046 PCT/US92/07022

These compounds are disclosed to be antagonists of AII which are useful for treating cardiovascular disorders such as hypertension.

F. Ostermeyer et al., in EP 475,898 (published
March 18, 1992) disclose heterocyclic N-substituted
derivatives of formula

These compounds are disclosed to be antagonists of AII which are useful for treating cardiovascular disorders such as hypertension.

P. Herold and P. Bühlmayer in EP 0 407 342 A2 disclose substituted pyrimidinones, pyrimidinethiones, and pyrimidinimines of the formula:

10

20

$$R_1$$
 R_2
 R_3
 R_4
 R_4

These are claimed to be antagonists of AII which are useful for treating hypertension.

E. Allen, et al. in EP 0 419 048 A (filed August 21, 1990) disclose a similar series of pyrimidinones which are claimed to be antagonists of AII

useful for the treatment of CHF and elevated intraocular pressure.

SUMMARY OF THE INVENTION

5 The present invention provides novel angiotensin II receptor antagonists of formula (I), pharmaceutical compositions containing compounds of formula (I) and therapeutic methods using them

10

wherein:

R1 is other than in the ortho position and is:

15 R^2 is

- (a) H
- (b) halo (F, Cl, Br, I),
- (c) C₁-C₄ alkyl,
- (d) C₁-C₄ alkoxy,
- (e) C₁-C₄ acyloxy,
- (c) of of actions
- (f) C_1-C_4 alkylthio,
- (g) C1-C4 alkylsulfinyl,
- (h) C₁-C₄ alkylsulfonyl,
- (i) hydroxy (C₁-C₄) alkyl,
- 25 (j) aryl (C₁-C₄) alkyl,

WO 93/04046 PCT/US92/07022

```
(k)
                -CO<sub>2</sub>H,
          (1)
                -CN,
                -CONHOR12,
          (m)
                -SO2NHR21,
          (n)
 5
          (o)
                -NH2,
                C1-C4 alkylamino,
          (p)
          (q)
                C1-C4 dialkylamino,
                -NHSO2R20,
          (r)
          (s)
                -NO2,
10
          (t)
                furyl,
          (u)
                arvl;
         wherein aryl is phenyl optionally substituted with
     one or two substituents selected from the group
     consisting of halo, C_1-C_4 alkyl, C_1-C_4 alkoxy, -NO_2,
     -CF3, C1-C4 alkylthio, -OH, -NH2, C1-C4 alkylamino, C1-C4
15
     dialkylamino, -CN, -CO2H, -CO2CH3, -CO2CH2CH3, -CO2-
     benzyl;
     R3 is
          (a)
                Η,
20
               halo,
          (b)
               C1-C4 alkyl,
         (c)
         (d)
                C1-C4 alkoxy,
         (e)
                C1-C4 alkoxyalkyl;
     R4 is
25
         (a)
                -CN,
         (b)
                -NO2,
         (c)
                -CO2R11;
     R<sup>5</sup> is
         (a)
                Η,
                C1-C6 alkyl,
30
         (b)
         (c)
                C3-C6 cycloalkyl,
         (d)
                C2-C4 alkenyl,
                C2-C4 alkynyl;
         (e)
```

PCT/US92/07022

WO 93/04046

```
R6 is
```

- (a) C_1-C_{10} alkyl,
- (b) C3-C8 alkenyl,
- (c) Ca-Ca alkvnvl,
- (d) C3-C8 cycloalkyl, 5
 - C4-C8 cycloalkenyl, (e)
 - (f) C4-C10 cycloalkylalkyl,
 - C5-C10 cycloalkylalkenyl, (q)
 - (h) C5-C10 cycloalkylalkynyl,
- 10 (i) - (CH₂) sZ (CH₂) mR⁵,
 - phenyl, optionally substituted with 1-2 (i) substituents selected from the group of halo, C1-C4 alkyl, C1-C4 alkoxy, -NO2, -NH2, -OH and benzyloxy,
 - benzyl, optionally substituted on the phenyl
- 15 ring with 1-2 substituents selected from the group of halo, C1-C4 alkyl, C1-C4 alkoxy and -NO2;
 - R^7 , R^8 , R^9 , and R^{10} are independently chosen from
 - (a) Η.
- 20 (b) C1-C8 alkyl,
 - (c) C1-C8 perfluoroalkyl,
 - (d) C3-C6 cycloalkyl,
 - (e) -NO2.
 - (f) -CN.
- -CONR15R16, 25 (g)
 - (h) -CO2R17,

 - (i) -OR18.
 - (i) -(CH₂)_nCONR¹⁵R¹⁶ where n is 1-4,
 - -(CH₂)_nCO₂R¹⁷ where n is 1-4, (k)
- 30 (1) (CH₂)_nOR¹⁸ where n is 1-4,
 - (m) aryl, wherein aryl is as defined above,
 - -CH2 arvl, wherein arvl is as defined above, (n)
 - R^7 and R^8 taken together are -(CH₂)+-, or (0)
 - (CH2) mX (CH2) a-,

```
R^9 and R^{10} taken together can be S, or O;
         (p)
     R^{11} is
         (a)
                Н,
                C1-C4 alkyl,
         (b)
                C1-C4 cycloalkyl,
 5
         (c)
         (d)
                phenyl,
                benzyl;
         (e)
     R12 is
         (a)
                Η,
10
         (b)
                methyl,
                benzyl;
         (c)
     R13 is
                -CH2CO2H,
         (a)
         (b)
                -C (CF3) 2OH,
                -CONHNHSO2CF3,
15
         (c)
                -CONHOR12,
         (d)
                -CONHSO2R20,
         (e)
                -CONHSO2NHR19,
         (f)
                -C (OH) R19PO3H2,
         (g)
                -NHCONHSO2R20,
         (h)
20
         (i)
                -NHPO3H2,
                -SO2NHCOR19,
         (i)
                -OPO3H2,
         (k)
                -OSO3H,
         (1)
                -PO (OH) R19,
25
         (m)
                -PO3H2,
         (n)
                -SO3H,
         (0)
                -SO2NHR19,
         (p)
                -NHSO2NHCOR19,
         (p)
                -SO2NHCONHR19,
30
         (r)
         (s)
```

(u) N=N. NH R4

5 (v)

 R^{14} is

- (a) H,
- (b) C₁-C₄ alkyl,
- 10 (c) aryl, wherein aryl is defined as above,
 - (d) benzyl,
 - (e) COR¹¹,
 - (f) CONHR¹¹;

R15 and R16 are independently

- 15 (a) H,
 - (b) C₁-C₆ alkyl,
 - (c) aryl, wherein aryl is as defined above,
 - (d) aryl (C_1 - C_4) alkyl, where aryl is as defined

above,

- 20 or taken together constitute a
 - (e) piperidine ring,
 - (f) morpholine ring,
 - (g) piperazine ring, optionally N-substituted with

C1-C6 alkyl, phenyl or benzyl;

 $25 R^{17} is$

- (a) H,
- (b) C₁-C₆ alkyl,
- (c) phenyl,
 - (d) benzyl;

 R^{18} is

- (a) H,
- (b) C_1-C_6 alkyl,
- (c) phenyl,
- 5 (d) benzvl;

 R^{19} is

10

15

- (a) H,
- (b) C₁-C₅ alkyl optionally substituted with a

substituent selected from the group consisting of aryl, where aryl is as defined above, heteroaryl where heteroaryl is as defined above, -OH, -SH, C_1 - C_4 alkyl, C_1 - C_4 alkoxy, C_1 - C_4 alkylthio, -CF3, halo, -NO2, -CO2H, -CO2CH3, -CO2-benzyl, -NH2, C_1 - C_4 alkylamino, C_1 - C_4

- dialkylamino, -PO3H2,
 - (c) aryl, where aryl is defined as above,
 - (d) -CH₂ aryl, where aryl is defined as above,
 - (e) heteroaryl,

wherein heteroaryl is an unsubstituted,

- monosubstituted or disubstituted 5- or 6-membered

 aromatic ring which can optionally contain from 1 to 3
 heteroatoms selected from the group consisting of O, N,
 and S and wherein the substituents are members selected
 from the group consisting of -OH, -SH, C1-C4 alkyl, C1-C4
 alkoxy, -CF3, halo, -NO2, -CO2H, -CO2CH3, -CO2-benzyl,
- 25 -NH₂, C₁-C₄ alkylamino, C₁-C₄ dialkylamino; R²⁰ is
 - (a) aryl, where aryl is defined as above,
 - (b) C₃-C₇ cycloalkyl,
 - (c) C₁-C₄ perfluoroalkyl,
- 30 (d) C₁-C₄ alkyl optionally substituted with a substituent selected from the group consisting of aryl, where aryl is as defined above, heteroaryl where

where aryl is as defined above, -OH, -SH, C1-C4 alkyl, C1-C4 alkoxy, C1-C4 alkylthio, -CF3, halo, -NO2, -CO2H,

WO 93/04046 PCT/US92/07022

.

15

20

25

11

```
-CO<sub>2</sub>CH<sub>3</sub>, -CO<sub>2</sub>-benzyl, -NH<sub>2</sub>, C_1-C<sub>4</sub> alkylamino, C_1-C<sub>4</sub>
      dialkylamino, -PO3H2,
                  heteroaryl, where heteroaryl is as defined
      above;
  5 R<sup>21</sup> is
           (a)
                н,
           (b) C<sub>1</sub>-C<sub>6</sub> alkyl,
          (c) phenyl,
           (d)
                benzyl,
10
    or taken together constitute a
          (e) piperidine ring,
          (f)
               morpholine ring,
                  piperazine ring, optionally N-substituted with
          (g)
          C1-C6 alkyl, phenyl or benzyl;
     X is
          (a) S,
          (b) 0,
          (c) SO,
          (d) SO2,
          (e) CHR14,
          (f) NR14;
     Z is
         (a) -O-,
          (b) -S-,
          (c) -NR11-;
     m is 1 to 5;
     n is 1 to 4;
30 q is 1 to 5;
     t is 2 to 5;
```

or a pharmaceutically acceptable salt thereof.

Preferred compounds of this invention are those of formula (I) wherein

R1 is in the para position and is

5

15



R6 is

- (a) C1-C10 alkyl,
- 10 (b) C₃-C₁₀ alkenyl,
 - (c) C₃-C₁₀ alkynyl,
 - (d) C3-C8 cycloalkyl,
 - (e) phenyl, optionally substituted with 1-2 substituents selected from the group of halo, C_1 - C_4 alkyl, C_1 - C_4 alkoxy, -NO₂, -NH₂, -OH and benzyloxy,
 - (f) benzyl, optionally substituted on the phenyl ring with one or two substituents selected from the group consisting of halo, C_1 - C_4 alkyl, C_1 - C_4 alkoxy and -NO₂;
- 20 R7, R8, R9, R10 are independently
 - (a) H,
 - (b) C₁-C₄ alkyl,
 - (c) C1-C4 perfluoroalkyl,
 - (d) C₃-C₆ cycloalkyl,
- 25 (e) phenyl, optionally substituted with one or two substituents selected from the group of halo, C₁-C₃ alkyl, C₁-C₃ alkoxy, NO₂, CF₃, NH₂, and OH, (f) R⁷ and R⁸ taken together are -(CH₂)_t-, or -(CH₂)_mX(CH₂)_σ-,
- 30 (g) R⁹ and R¹⁰ taken together can be S. O:

WO 93/04046 PCT/US92/07022

13

 R^{13} is

(a) -CONHSO2R20,

(b) -NHCONHSO2R20,

(c) -NHSO2NHCOR19,

(d) -PO3H2, 5

(e) -SO3H,

(f) -SO2NHR19,

(g) -SO2NHCOR19,

(h) -SO2NHCONHR19,

10 (i)

-CONH \(\frac{1}{N} \), \(\frac{1}{N} \). \(\frac{1}{N} \).

or a pharmaceutically acceptable salt thereof.

Still more preferred are compounds of the above 15 preferred scope formula (I) wherein

R6 is

(a) C1-C7 alkyl,

(b) C3-C4 alkenyl,

(c) C3-C4 alkynyl,

20 (d) phenyl, optionally substituted with 1-2 substituents selected from the group of halo, C_1-C_4

alkyl, C1-C4 alkoxy, -NO2, -NH2, -OH and benzyloxy;

R13 is

25

30

(a) -CONHSO2R20,

(b) -NHCONHSO2R20,

(c) -NHSO2NHCOR19,

(d) -SO2NHR19,

(e) $-SO_2NHCOR^{19}$, (f) -SO2NHCONHR19;

or a pharmaceutically acceptable salt thereof.

Most preferred due to their activity as angiotensin II antagonists are compounds of the more preferred scope wherein

R¹ is

or a pharmaceutically acceptable salt thereof.

Illustrative of the most preferred compounds of the invention are the following:

10

5

- N-((4'-(((4-oxo-2-propyl-8-thia-1,3-diazaspiro-((4.5))dec-1-en-3-yl-methyl))-((1,1'-biphenyl))-2ylsulfonyl))-benzamide
- 15 N-((4'-(((4-oxo-2-propyl-1,3-diazaspiro((4.4))non1-en-3-yl)methyl))((1,1'-biphenyl-2-ylsulfonyl))benzamide
- N-((4'-(((4-oxo-2-butyl-1,3-diazaspiro((4.4))non-1-20 en-3-yl)methyl))((1,1'-biphenyl-2-ylsulfonyl))benzamide
 - N-((4'-(4,5-dihydro-4,4-dimethyl-5-oxo-2-propyl-1H-imidazol-1-ylmethyl)((1,1'-biphenyl))-2-ylsulfonyl))-benzamide
- N-((4'-(4,5-dihydro-4,4-dimethyl-5-oxo-2-propyl-1H-imidazol-1-vlmethyl)((1,1'-biphenyl))-2-

ylsulfonyl))-4-chlorobenzamide

30

- N-((4'-(((4-oxo-2-propyl-8-thia-1,3-diazaspiro-((4.5))dec-1-en-3y-1-methyl))-((1,1'-biphenyl))-2vlsulfonvl))-hexanamide
- 5 • N-((4'-(((4-oxo-2-propyl-1,3-diazaspiro((2.4))hept-1-en-3-yl)methyl))((1,1'-biphenyl-2-ylsulfonyl))hexanamide
- N-((4'-(((4-oxo-2-propyl-1,3-diazaspiro((4.4))non-10 1-en-3-y1) methyl)) ((1,1'-biphenyl-2-y1sulfonyl))hexanamide
 - N-((4'-(((4-oxo-2-propyl-1,3-diazaspiro((4.5))dec-1en-3-yl)methyl))((1,1'-biphenyl-2-ylsulfonyl))hexanamide
- 15
 - N-((4'-(((4-oxo-2-butyl-1,3-diazaspiro((4.4))non-1en-3-yl) methyl)) ((1,1'-biphenyl-2-ylsulfonyl))hexanamide

- N-((4'-(4,5-dihydro-4,4-dimethyl-5-oxo-2-propyl-1Himidazol-1-ylmethyl) ((1,1'-biphenyl))-2ylsulfonyl))-hexanamide
- 25 N-((4'-(4,5-dihydro-4,4-dicyclopropyl-5-oxo-2propyl-1H-imidazol-1-ylmethyl) ((1,1'-biphenyl))-2ylsulfonyl))-hexanamide
- N-((4'-(4,5-dihydro-4,4-bistrifluoromethyl-5-oxo-2-30 propyl-1H-imidazol-1-ylmethyl) ((1,1'-biphenyl))-2ylsulfonyl))-hexanamide

- N-((4'-(4,5-dihydro-4,4-dimethyl-5-oxo-2-propyl-1Himidazol-1-ylmethyl)-3'-methyl((1,1'-biphenyl))-2vlsulfonvl))-hexanamide
- N-((4'-(4.5-dihydro-4.4-dimethyl-5-oxo-2-propyl-1Himidazol-1-ylmethyl)-4-propyl((1,1'-biphenyl))-2vlsulfonvl))-hexanamide
- N-((4'-(4,5-dihydro-4,4-dimethyl-5-oxo-2-propyl-1Himidazol-1-vlmethyl) ((1,1'-biphenyl))-2-10 vlsulfonyl))-trifluoroacetamide
- 3.5-dihvdro-5.5-dimethvl-2-propvl-3-[(2'-(N-((phenylsulfonyl)carboxamido)biphen-4-yl)methyl]-4Himidazol-4-one 15

Pharmaceutically suitable salts include both the metallic (inorganic) salts and organic salts; a list of which is given in Remington's Pharmaceutical Sciences, 17th Edition, page 1418 (1985). It is well known to one 20 skilled in the art that an appropriate salt form is chosen based on physical and chemical stability, flowability, hydroscopicity, and solubility. Preferred salts of this invention for reasons cited above include 25 potassium, sodium, calcium, and ammonium salts.

Detailed Description

Synthesis

30

The compounds of formula (I) may be prepared using the reactions and techniques described in this section. The reactions are performed in solvent suitable to the reagents and materials employed and suitable for the transformation being effected. It is understood by those skilled in the art of organic synthesis that the

functionality present on the imidazole and other portions of the molecule must be consistent with the chemical transformations proposed. This will frequently necessitate judgment as to the order of synthetic steps,

- 5 protecting groups required, deprotection conditions and activation of a benzylic position to enable attachment to nitrogen on the imidazole nucleus. Throughout the following section, not all compounds of formula (I) falling into a given class may necessarily be prepared
- by all the methods described for that class. Substituents on the starting materials may be incompatible with some of the reaction conditions required in some of the methods described. Such restrictions to the substituents which are compatible
- with the reaction conditions will be readily apparent to one skilled in the art and alternative methods described must then be used. The compounds of this application that have a chiral center may be resolved into the pure or partially pure optical isomers by any of the
- 20 appropriate procedures known to those skilled in the art.

The compounds of formula (I) can be prepared by alkylating the alkali-metal salt of the imidazoline 1a using appropriately protected benzyl halide, mesylate

25 (OMs), or tosylate (OTs) derivatives 2 as shown in Scheme 1

15

20

25

SCHEME 1

Q = I. Br. OMs. OTs

The alkylation produces a mixture of the two regioisomers using ether sodium hydride or potassium carbonate as base. The N^1 regionsomer is the major and the N3 is the minor products. These two isomers can be separated and purified using conventional separation 10 techniques such as chromatography or crystallization. In those cases where separation of regioisomers is difficult by conventional techniques, the mixture can be transformed into suitable derivatives that can be separated by usual separation methods. They possess distinct physical and biological properties.

The benzyl halides of formula 2 can be prepared as described in European Patent Applications 400 974; 401 030; 400,835 and references therein.

The starting imidazolinones are readily available by any number of standard methods. For example imidazolinone of formula 1 can be prepared as shown in Scheme 2. The amino nitrile 3 is readily obtainable from aldehydes and ketones via the Strecker Synthesis and various modifications thereof ($R^7 = R^8 = CF_3$, Y. V. Zeifman, N. P. Gambaryan, I. L. Knunyants, Dokl. Acad.

Nauk.S.S.S.R., 153, 1334, 1963). Treatment of the amino nitrile with triethyl amine and one equivalent of the appropriate acyl or aroyl chloride 4 in methylene chloride at room temperature overnight, gives the 5 corresponding amidonitrile 5. Alternatively, the nitrile can be made following the procedure described in German patent disclosure DE3704100Al. The nitrile can be hydrolyzed to the diamide 6 using standard procedures such as treatment with hydrochloric acid followed by ammonium hydroxide. Treatment of the diamide with 1 N sodium hydroxide as described in E. Mohr, J. Pract. Chem., 81, 49, 1910, gives the imidazolinone 1.

Alternatively, imidazolinones of formula 1 can also be prepared as shown in Scheme 3. Treatment of the amino acid 7 with tert-butyl pyrocarbonate 4 with two or 15 more equivalents of base gives the BOC (tertbutyloxycarbonyl) protected amino acid 5, M. Bodanszky and A. Bodanszky, The Practice of Peptide Chemistry, 1984. The protected amino amide 6 can be synthesized 20 from the active ester followed by treatment with ammonia. Deprotection using HCl gas gives the amino amide hydrochloride 7. Treatment with two or more equivalents of base and the appropriate acyl or aroyl

SCHEME 2

5 chloride gives the diamide 6 which can be cyclized by treatment with 1 N sodium hydroxide as described above.

Likewise, compound 6 may be obtained by reacting amino acid with the requisite acid chloride by either a Schotten-Baumann procedure, or simply stirring in a solvent such as methylene chloride in the presence of base such as sodium bicarbonate, pyridine or triethyl amine followed by coupling reaction with ammonia via a variety of amide or peptide forming reactions such as DCC coupling, azide coupling, mixed anhydride synthesis or any other coupling procedure familiar to one skilled in the art.

The use of 1-amino-1-cycloalkylcarboxylic acids in the above procedure provides the imidazolinone starting materials for the preparation of the spiro-substituted imidazolinones of formula (I).

SCHEME 3

- 5 Imidazolinones of formula 1 can also be prepared following the procedure described in Japanese Patent disclosure JP 58055467, and the procedure by H. Lehr, J. Am. Chem. Soc., 75, 3640, 1953 and references therein.
- Imidazolinones of formula 1 wherein R7 an R8 are both phenyl can be prepared as shown in Scheme 4 by

20

reaction of benzil 12 with alkyl or aryl amidine hydrochloride 13, A. W. Cox, Org. Syn., 1, 5, R. T. Boere, R. T. Oakley, R. W. Reed, J. Organomet. Chem., 331, 161, 1987, in the presence of base such as 1 N sodium hydroxide, G. Rio and A. Rajon, Bull. Soc. Chim. France, 543, 1958 and references therein.

SCHEME 4

The imidazoline thiones of formula 15 can be prepared by treatment of the requisite alkylated imidazolinone 14 with Lawesson's reagent or phosphorus pentasulfide as described in M. P. Cava and M. I. Levinson, Tetrahedron, 41, 5061, 1985 (Scheme 5).

SCHEME 5

The compounds of this invention and their preparation can be understood further by the following

WO 93/04046 PCT/US92/07022

23

examples which do not constitute a limitation of the invention. In these examples, unless otherwise indicated, all temperatures are in degrees centigrade and parts and percentages are by weight.

5

EXAMPLE 1

N=((4'-(((4-oxo-2-propyl-8-thia-1,3-diazaspiro-) ((4.5))dec-1-en-3-yl-methyl))-((1,1'-biphenyl))-2ylsulfonyl))-benzamide

PART A: Preparation of 4-Amino-4-cyanotetrahydrothiopyrane

15

20

25

Sodium cyanide (2.11 g, 43 mmol) was dissolved in water (40 ml). Ammonium chloride (2.53 g, 47.3 mmol) was added followed by a solution of tetrahydrothiopyran-4-one (5.0 g, 43 mmol) in methanol (40 ml). The mixture was stirred at room temperature under N₂ overnight. The mixture was diluted with H₂O and extracted with methylene chloride. The combined organic layer was washed with brine, dried over MgSO₄ and concentrated. The residue was chromatographed on silica gel eluting with hexane-ethyl acetate (1:1) to give 5.28 g of white solid (86%). MS m/e 143.0 (M+H)+; 1HMMR (CDC13/TMS) δ 1.60-2.00 (m, 4H, CH₂), 2.25(d, 2H, CH₂), 2.62-3.00 (m, 4H, CH₂ and NH₂); IR(KBr, cm⁻¹) 2218.6(s, CN), 3371.3 & 3302.5(d, NHo).

PART B: Preparation of 4-N-butyramido-4-cyanotetrahydrothiopyrane

Butyryl chloride (5.8 ml, 40.8 mmol) was added

dropwise to a cooled mixture of 4-Amino-4-cyanotetrahydrothiopyrane (5.28 g, 37.1 mmol) and triethyl
amine (5 ml) in methylene chloride (150 ml). The
mixture was stirred for 3 hours at room temperature
after which it was poured into 1N Hcl (50 ml). The

organic layer was washed with 1N Hcl (2x50 ml), 1N NaOH
(2x50 ml), dried (MgSO₄) and concentrated. The residue
was triturated with hexane to give a white solid (7.50
g, 95%). MS m/e 213, (M+H)+, 1HNMR (CDCl₃/TMS) & 0.98(t,
3H, CH₃), 1.68(m, 2H, CH₂), 1.96(m, 2H, CH₂), 2.20(t, 2H,
NH).

PART C: Preparation of 4-Aminocarbonyl-4-N-butyramidotetrahydrothiopyrane

20

25

30

4-N-butyramido-4-cyanotetrahydro-thiopyrane (7.5 g, 35.3 mmol) was dissolved in concentrated hydrochloric acid (50 ml) at 0°C. Cold water (175 ml) was added immediately followed by treatment with concentrated ammonium hydroxide to pH 5-6. The mixture was extracted successively with methylene chloride. The organic layer was combined, washed with brine, dried over MgSO₄ and concentrated to give white solid (7.0 g, 86%). MS m/e 231 (M+H)+, 1HNMR (DMSO-d6/TMS) δ 0.86(t, 3H, CH₃), 1.50(s, 2H, CH₂), 1.90(t, 2H, CH₂), 2.18(t, 2H, CH₂), 2.20-2.47(m, 4H, CH₂), 2.76(t, 2H, CH₂), 6.90(t, 2H, NH₂), 7.68(s, 1H, NH).

PART D: Preparation of 4-0xo-2-propyl-8-thia-1,3-diazaspiro-(4,5)dec-1-ene

4-Aminocarbonyl-4-N-butyramido-tetrahydrothiopyrane

(7.0 g, 30.4 mmol) was heated with 1N sodium hydroxide

(50 ml) for 30 minutes. The mixture was cooled to room temperature and some solid material was filtered off.

The filtrate was neutralized with aqueous HCl and the white precipitate formed was filtered and dried (2.63

g). The aqueous layer was extracted with methylene chloride. The combined organic layer was washed with brine, dried and concentrated to a white solid (1.35 g). A total of 3.98 g of product was isolated (62%). MS m/e 213 (M+H)+, lHNMR (CDCl3/TMS) & 1.00(t, 3H, CH3), 1.60
15 1.80 (m, 4H, CH2), 2.05 (m, 2H, CH2), 2.44(t, 2H, CH2), 2.75 (m, 2H, CH2), 3.02(t, 2H, CH2), 8.30(s, 1H, NH).

PART E: Preparation of N-[(4'-[((4-oxo-2-propyl-8-thia-1.3-diazaspiro-[(4.5)]dec-1-en-3-yl-methyl)]-N-tertbutyl[(1.1'-biphenyl)]-2-sulfonamid

4-0xo-2-propyl-8-thia-1,3-diazaspiro-(4,5)dec-1-ene
(0.83 g, 3.9 mmol) was dissolved in dimethyl formamide
(20 ml). Sodium hydride (0.18 g of 80% dispersion in
25 mineral oil) was added portionwise. The mixture was
allowed to stir at room temperature for 15 minutes. 4'bromomethyl-N-tert-butyl(1,1'-biphenyl)-2-sulfonamide
(1.5 g, 3.9 mmol) was added. The mixture was stirred at
room temperature under N₂ for 24 h. The solvent was
30 removed in vacuo. The residue was dissolved in ethyl
acetate and washes with water and brine. It was then
dried over MgSO₄, concentrated and chromatographed on
silica gel eluting with hexane-ethyl acetate (1:1) to
yield 1.65 g of white solid (82%). MS m/e 514.2 (M+H)+.

15

20

25

30

1HNMR(CDC1₃/TMS) δ 0.98(t, 3H, CH₃), 0.99(s, 9H, CH₃),
1.74(m, 4H, CH₂), 2.10(m, 2H, CH₂), 2.34(t, 2H, CH₂),
2.79(m, 2H, CH₂), 3.09(t, 2H, CH₂), 3.45(s, 1H, NH),
4.73(s, 2H, CH₂Ar), 7.25(m, 3H, ArH), 7.50(m, 4H, ArH),
8.18(d, 1H, ArH).

PART F: Preparation of N-((4'-((4-oxo-2-propyl-8-thia-1,3-diazaspiro-((4.5))dec-1-en-3-yl-methyl))((1,1'biphenyl))-2-sulfonamide

N-((4'-(((4-oxo-2-propyl-8-thia-1,3-diazaspiro-((4.5))dec-1-en-3-yl-methyl))-N-tert-butyl((1,1'-biphenyl))-2-sulfonamide (1.56 g, 3.0 mmol) was refluxed with trifluoroacetic acid (10 ml) under N₂ for 2 h. The solvent was removed in vacuo and the residue was dissolved in methylene chloride. The organic solution was washed with aqueous NaHCO3 and brine. It was filtered through phase transfer paper and concentrated to an off-white solid (1.31 g, 95%). MS m/e 458.0 (M+H)+, 1 HNMR(CDC13/7MS) δ 0.97(t, 3H, CH₃), 1.60-1.82 (m, 4H, CH₂), 2.10 (m, 2H, CH₂), 2.36(t, 2H, CH₂), 2.78 (m, 2H, CH₂), 3.09(t, 2H, CH₂), 4.21(s, 2H, NH₂), 4.73 (s, 2H, CH₂Ar), 7.20 (d, 2H, ArH), 7.30 (d, 1H, ArH), 7.42-7.63 (m, 4H, ArH), 8.17 (d, 1H, ArH), 7.

PART G: Preparation of N-((4'-(((4-oxo-2-propyl-8-thia-1.3-diazaspiro-((4.5))dec-1-en-3-yl-methyl))-((1.1'biphenyl))-2-ylsulfonyl))-benzamide

1,1'-Carbonyl diimidazole (1.26 g, 7.8 mmol) and benzoic acid (0.96 g, 7.8 mmol) was refluxed with tetrahydrofurane (30 ml) under N_2 for 2 h. The mixture was cooled to room temperature, and a solution of N-((4'-(($(4-\infty x)^2-propyl-8-thia-1,3-diazaspiro-$

((4.5))dec-1-en-3-vl-methyl))((1,1'-biphenyl))-2sulfonamide (1.20 g, 2.6 mmol) and 1,8diazabicvclo[5.4.0]undec-7-ene (1.2 ml, 7.8 mmol) in THF (30 ml) was added. The reaction mixture was then 5 refluxed for 1.5 h. The mixture was cooled and poured into 20 ml of 25% aqueous citric acid. It was extracted with ethyl acetate. The combined organic solution was washed with brine, dried over MgSO4 and concentrated. the residue was chromatographed on silica gel eluting 10 with 5% methanol in methylene chloride to yield 1.3 g of the desired product (91%). M.P. 105-108°C, MS m/e 561 (M+H)+, 1 HNMR(CDCl₃/TMS) δ 0.98(t, 3H, CH₃), 1.60-1.80(m, 4H, CH₂), 2.08(t, 2H, CH₂), 2.34(t, 2H, CH₂), 2.78(m, 2H, CH2), 3.08(t, 2H, CH2), 4.65(s, 2H, CH2Ar), 6.99(d, 2H, 15 ArH), 7.23(d, 1H, ArH), 7.42(m, 4H, ArH), 7.60(m, 4H, ArH), 8.40(d, 1H, ArH).

EXAMPLE 2

N = ((4' - (((4-oxo-2-propvl-1, 3-diazaspiro((4,4))non-1-en-3-v1)methv1))((1,1'-biphenv1-2-v1sulfonv1))-benzamide 20

PART A: Preparation of 4-0xo-2-propvl-1.3-diazaspiro-(4,4)non-1-ene

25 1-Amino-1-cyclopentane carboxylic acid methyl ester (10.1 g, 70.6 mmol), ethyl butanimidate hydrochloride (12.7 g, 84.7 mmol) and triethyl amine (17 ml) was refluxed in benzene (50 ml) under No overnight. The solvent was removed in vacuo and the residue was dissolved in methylene chloride, washed with water and 30 brine, and concentrated. The crude product mixture was chromatographed on silica gel eluting with ethyl acetate to give 8.47 g of colorless oil (67%). MS m/e 181.1

•

10

15

 $(M+H)^+$, 1 HNMR(CDCl₃/TMS) δ 1.00(t, 3H, CH₃), 1.50-2.20(m, 10H, CH₂), 2.41(t, 2H, CH₂), 9.09(br.s, 1H, NH).

PART B: N-((4'-(((4-oxo-2-propyl-1,3-diazaspiro5 ((4.4))non-1-en-3-yl)methyl))((1,1'-biphenyl-2vlsulfonyl))-bebzamide

The titled compound was prepared from 4-0xo-2-propyl-1,3-diazaspiro-(4,4)non-1-ene and 4'-bromomethyl-N-tert-butyl(1,1'-biphenyl)-2-sulfonamide according to the procedures described in Part E, F and G of Example 1. M.P. 118-125°C, MS m/e 529 (M+H)+, 1HNMR (DMSO-d6/TMS) & 0.89(t, 3H, CH3), 1.50-1.92(m, 10H, CH2), 2.34(t, 2H, CH2), 4.68(s, 2H, CH2Ar), 7.00(d, 2H, ArH), 7.10-7.62(m, 10H, ArH), 8.10(d, 1H, ArH).

Compounds 1-630 in Table 1 can be prepared by the procedures described in Examples 1 and 2 employing the appropriately substituted imidazolines and benzyl halides or mesulates.

TABLE 1

5.	EX.	R6	R ⁷	R ⁸	R ⁹ R ¹⁰	R ¹³	R ² , R ³	MS (M+H)+
	1	n-Pr	- (CH ₂	2S (CH) 2-	0	-so ₂ nHCoC ₆ H ₅	Н,Н	561
	2	n-Pr	- (CH ₂	4-	0	-so ₂ NHCOC ₆ H ₅	н, н	529
	3	n-Pr	- (CH ₂) 4-	0	$-so_2$ NHCO (n-C ₄ H ₉)	Н,Н	
10	4	n-Bu	- (CH ₂) 4-	0	-so ₂ NHCOC ₆ H ₅	Н,Н	543
	5	n-Pr	CH ₃	СНЗ	0	-so2NHCOC6H5	Н, Н	504
	6	n-Pr	CH ₃	СНЗ	0	-CONHSO2CH2C6H5	н, н	
	7	n-Pr	CH ₃	CH ₃	0	-so ₂ NHCO (n-C ₅ H ₁₁)	к Н,Н	
	8	n-Pr	CH ₃	CH ₃	0	$-so_2$ NHCO(cy- C_3H_5)	н,н	
15	9	n-Pr	CH3	CH ₃	0	-so ₂ NHCOCH ₂ C ₆ H ₅	н,н	
	10	n-Pr	CH3	CH ₃	0	-CH2CO2H	н, н	
	11	n-Pr	CH3	CH3	0	-C (CF ₃) 2OH	н, н	
	12	n-Pr	CH3	CH3	0	-CONHNHSO2CF3	н, н	
	13	n-Pr	CH3	CH3	0	-CONHOCH3	н, н	
20	14	n-Pr	CH3	CH ₃	0	-conhso ₂ c ₆ H ₅	н, н	
	15	n-Pr	CH3	CH3	0	-P03H2	н, н	
	16	n-Pr	CH3	CH3	0	-so ₂ NHCOC ₆ H ₅	CH3,	H
	17	n-Pr	снз	CH ₃	0	$-so_2$ NHCO (n-C ₅ H ₁₁)	CH ₃ ,	H
	18	n-Pr	СНЗ	CH ₃	0	$\hbox{-so}_2\hbox{\tt NHCO}(\hbox{\tt cy-C}_3\hbox{\tt H}_5)$	CH3,	H
25	19	n-Pr	CH3	CH ₃	0	$-so_2$ NHCOCH $_2$ C $_6$ H $_5$	CH3,	Н
	20	n-Pr	СНЗ	СНЗ	0	-so ₃ H	CH3,	Н
	21	n-Pr	CH ₃	CH3	0	-so2NHCONH (n-Bu)	CH3,	Н
	22	n-Pr	CH ₃	CH3	0	-so2NH(C5NH4)	CH3,	Н

	EX	. R6	R7	R ⁸	R9R10	R ¹³	\mathbb{R}^2 , \mathbb{R}^3
	23	n-Pr	CH ₃	сн3	0	$-so_2$ NHCONH (n-C ₅ H ₁₁)	сн3,н
	24	n-Pr	CH ₃	CH ₃	0	-NHSO ₂ NHCO (i- C_5H_{11})	CH ₃ ,H
	25	n,-Pr	снз	сн3	0	- $\mathrm{nHso}_2\mathrm{nHco}\left(\mathrm{cy-C}_3\mathrm{H}_5\right)$	CH ₃ ,H
5	26	n-Pr	CH ₃	сн3	0	-NHSO2NHCOCH2C6H5	CH ₃ ,H
	27	n-Pr	CH3	сн3	0	-so2nHCOC6H5	C1,H
	28	n-Pr	CH ₃	сн3	0	-so ₂ NHCO(n-C ₅ H ₁₁)	Cl,H
	29	n-Pr	CH ₃	сн3	0	$-so_2$ NHCO (cy- c_3H_5)	Cl,H
	30	n-Pr	CH ₃	сн3	0	-so ₂ NHCOCH ₂ C ₆ H ₅	C1,H
10	31	n-Pr	CH ₃	снз	0	-opo ₃ H ₂	Cl,H
	32	n-Pr	CH ₃	сн3	0	-PO (OH) CH2C6H5	Cl,H
	33	n-Pr	CH ₃	сн3	0	-oso ₃ H	Cl,H
	34	n-Pr	CH ₃	CH ₃	0	-NHPO3H2	Cl,H
	35	n-Pr	CH ₃	сн3	0	-so ₂ NH ₂	Cl,H
15	36	n-Pr	CH ₃	сн3	0	-so ₂ NHC ₂ H ₅	Cl,H
	37	n-Pr	CH ₃	CH ₃	0	-so ₂ NHC ₁₀ H ₇	Cl,H
	38	n-Pr	CH ₃	CH ₃	0	-so2nHCOC6H5	F,H
	39	n-Pr	CH ₃	CH ₃	0	-SO2NHCO (n-C5H11)	F,H
	40	n-Pr	CH ₃	CH ₃	0	-so2NHCO(cy-C3H5)	F,H
20	41	n-Pr	CH ₃	CH ₃	0	-so2nHCOCH2C6H5	F,H
	42	n-Pr	CH ₃	CH ₃	0	-NHSO2NHCO(n-Bu)	F,H
	43	n-Pr	CH ₃	CH ₃	0	-SO2NHCONH (n-Bu)	F,H
	44	n-Pr	CH ₃	CH ₃	0	-SO2NHCO(i-Bu)	F,H
	45	n-Pr	CH ₃	сн3	0	-so2NHCO(4-HOC6H5)	F,H
25	46	n-Pr	CH ₃	сн3	0	-so ₂ nHCOCH ₂ C ₆ H ₅	F,H
	47	n-Pr	CH ₃	СН3	0	-so2NHCO(4-C5NH4)	F,H
	48	n-Pr	CH ₃	СН3	0	-so ₂ nhConhCh ₂ C ₆ H ₅	F,H
	49	n-Pr	CH ₃	сн3	0	-so ₂ NHCOC ₆ H ₅	H,n-Pr
	50	n-Pr	CH ₃	сн3	0	-SO2NHCO(n-C5H11)	H,n-Pr
30	51	n-Pr	CH ₃	сн3	0	$-so_2$ NHCO(cy- c_3H_5)	H,n-Pr
	52	n-Pr	CH ₃	сн3	0	-so ₂ nHCOCH ₂ C ₆ H ₅	H,n-Pr
	53	n-Pr	CH ₃	CH ₃	0	-сн ₂ со ₂ н	H,n-Pr
	54	n-Pr	CH ₃	снз	0	-C (CF ₃) ₂ OH	H,n-Pr
	55	n-Pr	CH ₃	CH ₃	0	-CONHNHSO2CF3	H,n-Pr

		EX	. R ⁶	_R 7	R8	R9R10	R ¹³	R ² , R ³
		56	n-Pr	СНЗ	Сн3	О	-соннон	H,n-Pr
		57	n-Pr	CH ₃	СНЗ	О	-соиносн3	H, n-Pr
•		58	n-Pr	CH ₃	СН3	0	-соnносн ₂ с ₆ с ₅	H,n-Pr
	5	59	n-Pr	СНЗ	СНЗ	0	-so ₂ NHCOC ₆ H ₅	Cl,n-Pr
,		60	n-Pr	CH ₃	СН3	0	-SO2NHCO (n-C5H11)	Cl,n-Pr
		61	n-Pr	CH ₃	СН3	0	-so2NHCO(cy-C3H5)	F,n-Pr
		62	n-Pr	CH ₃	CH ₃	О	-so ₂ NHCOCH ₂ C ₆ H ₅	F,n-Pr
		63	n-Pr	CH3	CH ₃	0	-CONHSO2 (4-C1C6H4)	Cl,n-Pr
	10	64	n-Pr	CH ₃	CH ₃	0	-conhso ₂ c ₂ F ₅	F,n-Pr
		65	n-Pr	CH ₃	СНЗ	0	-conhso ₂ c ₂ h ₄ oh	Cl,n-Pr
		66	n-Pr	CH ₃	CH ₃	0	-conhso2c2H4co2H	F,n-Pr
		67	n-Pr	СНЗ	CH3	0	-conhso ₂ c ₂ H ₄ NH ₂	F,n-Pr
		68	n-Pr	CH ₃	СНЗ	0	-conhso ₂ nh ₂	Cl,n-Pr
	15	69	n-Bu	CH3	сн3	0	-so2NHCOC6H5	н,н
		70	n-Bu	CH ₃	СНЗ	0	-so ₂ NHCO(n-C ₅ H ₁₁)	н,н
		71	n-Bu	CH ₃	СН3	0	-so2nHCO(cy-C3H5)	H,H
		72	n-Bu	CH ₃	снз	0	-so2NHCOCH2C6H5	н, н
		73	n-Bu	CH ₃	снз	0	-conhso2nhc2h5	н, н
	20	74	n-Bu	CH ₃	снз	0	-conhso2nhc6h5	н, н
		75	n-Bu	CH ₃	CH ₃	0	-conhso2nhch2c6H5	н, н
		76	n-Bu	CH ₃	CH ₃	0	-conhso2nh(4-c5nh4)	н,н
		77	n-Bu	СН3	CH ₃	0	-C (OH) CH3PO3H2	н,н
		78	n-Bu	сн3	снз	0	-C (OH) HPO3H2	Н,Н
	25	79	n-Bu	сн3	CH ₃	0	-so ₂ nHcoc ₆ H ₅	сн3,н
		80	n-Bu	CH ₃	CH ₃	0	-so2NHCO(n-C5H11)	сн3,н
		81	n-Bu	CH ₃	СНЗ	0	-so2NHCO(cy-C3H5)	сн3,н
		82	n-Bu	CH ₃	CH3	0	-so2NHCOCH2C6H5	сн ₃ , н
		83	n-Bu	CH3	снз	0	-NHCONHSO2C2H5	СH3, Н
	30	84	n-Bu	сн3	СНЗ	0	-NHCONHSO ₂ (i-Bu)	Сн ₃ ,н
		85	n-Bu	Сн3	сн3	0	-NHCONHSO ₂ (n-C ₅ H ₁₁)	Сн ₃ , н
		86	n-Bu	CH3	СНЗ	0	-so ₂ nhconh (n-c ₅ H ₁₁)	сн ₃ ,н
		87	n-Bu	CH ₃	снз	0	$-so_2$ NHCO($i-c_5H_{11}$)	сн ₃ ,н
		88	n-Bu	СНЗ	CH3	0	-so2NHCO(cy-C3H5)	сн ₃ , н

•	EX.	R6	R ⁷	R8	R9R10	R13	\mathbb{R}^2 , \mathbb{R}^3
	89	n-Bu	СНЗ	сн3	О	-so ₂ NHCOCH ₂ C ₆ H ₅	СН3,Н
	90	n-Bu	сн3	сн3	О	-so ₂ NHCOC ₆ H ₅	Cl,H
	91	n-Bu	CH3	CH3	0	$-so_2$ NHCO (n-C ₅ H ₁₁)	Cl,H
5	92	n-Bu	СН3	CH ₃	0	-so2NHCO(cy-C3H5)	Cl,H
	93	n-Bu	CH ₃	CH ₃	0	-so ₂ nhcoch ₂ c ₆ H ₅	Cl,H
	94	n-Bu	CH ₃	CH ₃	0	-NHP03H2	Cl,H
	95	n-Bu	CH ₃	снз	0	$-\text{NHSO}_2\text{NHCO}\left(\text{Cy-C}_3\text{H}_5\right)$	Cl,H
	96	n-Bu	CH3	CH3	0	-SO2NHCONH(i-Bu)	Cl,H
10	97	n-Bu	CH ₃	сн3	0	-PO (OH) (n-C ₅ H ₁₁)	Cl,H
	98	n-Bu	CH ₃	CH ₃	0	-PO(OH)(1-C5H11)	Cl,H
	99	n-Bu	CH ₃	CH3	0	-PO (OH) C3H7	Cl,H
	100	n-Bu	CH3	CH3	0	-so2NHCH2C6H5	Cl,H
	101	n-Bu	CH ₃	CH ₃	0	-so2NHC6H5	F,H
15	102	n-Bu	CH ₃	CH ₃	0	-so ₂ NHCO(n-C ₅ H ₁₁)	F,H
	103	n-Bu	снз	CH ₃	0	-SO2NHCO(cy-C3H5)	F,H
	104	n-Bu	сн3	CH ₃	0	-so2NHCOCH2C6H5	F,H
	105	n-Bu	сн3	CH ₃	0	-SO2NH (n-Bu)	F,H
	106	n-Bu	CH ₃	CH ₃	0	-SO ₂ NH(i-Bu)	F,H
20	107	n-Bu	сн3	CH ₃	0	-SO2NHCONH(i-Bu)	F,H
	108	n-Bu	снз	CH ₃	0	-so ₂ NH (n-C ₅ H ₁₁)	F,H
	109	n-Bu	CH ₃	CH ₃	0	$-so_2$ NH(i-C ₅ H ₁₁)	F,H
	110	n-Bu	СНЗ	сн3	0	-NHSO2NHCO(cy-C3H5)	F,H
	111	n-Bu	CH3	CH ₃	0	-so ₂ NHCOCH ₂ C ₆ H ₅	F,H
25	112	n-Bu	CH3	CH3	0	-so ₂ NHCOC ₆ H ₅	H,n-Pr
	113	n-Bu	CH ₃	сн3	0	-so ₂ NHCO(n-C ₅ H ₁₁)	H,n-Pr
	114	n-Bu	CH ₃	CH ₃	0	$-so_2$ NHCO(cy-C ₃ H ₅)	H,n-Pr
	115	n-Bu	сн3	сн3	0	-so ₂ nhcoch ₂ c ₆ h ₅	H,n-Pr
	116	n-Bu	CH3	CH3	0	-SO2NHCO(n-Bu)	H, n-Pr
30	117	n-Bu	CH ₃	CH ₃	0	-SO2NHCONH (n-Bu)	H,n-Pr
	118	n-Bu	CH ₃	CH3	0	-SO2NHCONH(i-Bu)	H, n-Pr
	119	n-Bu	снз	сн3	0	$-so_2NH (n-c_5H_{11})$	H,n-Pr
	120	n-Bu	СНЗ	сн3	0	$-so_2$ NH(i-C ₅ H ₁₁)	H,n-Pr
	121	n-Bu	снз	сн3	0	$-so_2$ NH(cy- C_3 H ₅)	H,n-Pr

	EX	. R6	R ⁷	R8	R9R10	R13	R^2, R^3
	122	n-Bu	сн3	СНЗ	0	$-so_2$ NHCH $_2$ C $_6$ H $_5$	H,n-Pr
	123	n-Bu	сн3	CH ₃	0	-so2NHCOC6H5	Cl,n-Pr
	124	n-Bu	снз	CH ₃	0	-so ₂ NHCO (n-C ₅ H ₁₁)	Cl,n-Pr
5	125	n-Bu	снз	снз	0	$-so_2$ NHCO (cy- c_3H_5)	F,n-Pr
	126	n-Bu	снз	снз	0	-so2NHCOCH2C6H5	F,n-Pr
	127	n-Bu	снз	снз	0	-So ₂ NH (n-Bu)	Cl,n-Pr
	128	n-Bu	CH ₃	СНЗ	0	-so ₂ NH(i-Bu)	F,n-Pr
	129	n-Bu	CH ₃	сн3	0	-contso ₂ (n-c ₅ H ₁₁)	Cl,n-Pr
10	130	n-Bu	СНЗ	снз	0	$-conhso_2 (i-c_5H_{11})$	F,n-Pr
	131	n-Bu	CH ₃	СНЗ	0	-conhso ₂ (cy-c ₃ H ₅)	F,n-Pr
	132	n-Bu	CH3	CH3	0	-conhso2ch2c6h5	Cl,n-Pr
	133	Ph	СНЗ	СНЗ	0	-so2NHCOC6H5	н, н
	134	Ph	снз	CH3	0	-SO2NHCO(n-C5H11)	н, н
15	135	Ph	сн3	CH3	0	-SO2NHCO(cy-C3H5)	н, н
	136	p-F-Ph	сн3	CH ₃	0	-so2NHCOCH2C6H5	н, н
	137	p-F-Ph	СНЗ	CH3	0	-SO2NHCO (n-Bu)	н, н
	138	Ph	снз	СНЗ	0	-So ₂ NHCo(i-Bu)	н, н
	139	iPr	СНЗ	СНЗ	0	-so ₂ NHCO (n-C ₅ H ₁₁)	н,н
20	140	Ph	CH3	снз	0	$-so_2$ NHCO($i-c_5H_{11}$)	н, н
	141	Ph	СНЗ	CH3	0	-so2NHCO(cy-c3H5)	н, н
	142	Ph	CH3	снз	0	-so ₂ nhcoch ₂ c ₆ h ₅	н,н
	143	n-Pr	- (CH ₂) 4-	0	-conhso2ch2c6H2	н, н
	144	n-Pr	- (CH ₂) 4-	0	$-so_2$ NHCO(n-C ₅ H ₁₁)	H,H
25	145	n-Pr	-(CH ₂) 4-	0	-so2NHCO(cy-C3H5)	н, н
	146	n-Pr	-(CH ₂) ₄ -	0	-so2NHCOCH2C6H5	н, н
	147	n-Pr	- (CH ₂) 4-	0	-CH2CO2H	н,н
	148	n-Pr	- (CH ₂) ₄ -	0	-C (CF ₃) ₂ OH	н, н
	149	n-Pr	-(CH ₂) ₄ -	0	-conhnhso2cf3	Н,Н
30	150	n-Pr	-(CH ₂) ₄ -	0	-соиносн ₂ с ₆ н ₅	н,н
	151	n-Pr	- (CH ₂	4-	0	-соиносн3	н, н
	152	n-Pr	- (CH ₂	4-	0	-so2NHCOC6H5	сн ₃ ,н
	153	n-Pr	-(CH ₂	4-	0	$-so_2$ NHCO (n-C ₅ H ₁₁)	СH3, Н
	154	n-Pr	-(CH ₂	4-	0	$-so_2$ nhco (cy- c_3H_5)	СН3,Н

	EX.	. R6	R ⁷	R8	R ⁹ R	10 R ¹³	R ² ,	R3
	155	n-Pr	- (CH	2)4-	0	-so ₂ nhcoci	H ₂ C ₆ H ₅ СН ₃	, H
	156	n-Pr	- (CH	2)4-	0	-so ₂ nhco (1	n-Bu) CH ₃	3,Н
	157	n-Pr	- (CH	2)4-	0	-so ₂ nhconi	H(n-Bu) CH3	,Н
5	158	n-Pr	- (CH	2)4-	0	-CONHSO ₂ (L-Bu) CH ₃	, Н
	159	n-Pr	- (CH	2)4-	0	-conhso ₂ (r	n-C ₅ H ₁₁) СН ₃	, н
	160	n-Pr	- (CH	2)4-	0	-conhso ₂ (i	i-C ₅ H ₁₁) СН ₃	, н
	161	n-Pr	- (CH	2)4-	0	-conhso2 (су-С ₃ н ₅) Сн ₃	, н
	162	n-Pr	- (CH	2)4-	0	-conhso ₂ ci	₁₂ С ₆ н ₅ сн ₃	, Н
10	163	n-Pr	- (CH	2)4-	0	-so ₂ nhcoc	₅ H ₅ Cl,	H
	164	n-Pr	- (CH	2)4-	0	-so ₂ nhco(r	1-C ₅ H ₁₁) C1,	н
	165	n-Pr	- (CH	2)4-	0	-so2NHCO($cy-c_3H_5)$ cl,	н
	166	n-Pr	- (CH	2)4-	0	-so ₂ NHCOCH	1 ₂ C ₆ H ₅ C1,	H
	167	n-Pr	- (CH	2)4-	0	-conhso ₂ ni	I(n-Bu) Cl,	н
15	168	n-Pr	- (CH	2)4-	0	-SO2NHCONE	I(n-Bu) Cl,	Н
	169	n-Pr	- (CH	2)4-	0	-NHCONHSO ₂	(i-Bu) Cl,	H
	170	n-Pr	- (CH	2)4-	0	-NHCONHSO ₂	(n-C ₅ H ₁₁) Cl,	H
	171	n-Pr	- (CH	2)4-	0	-NHCONHSO2	(i-c ₅ H ₁₁) cl,	H
	172	n-Pr	- (CH ₂	2)4-	0	-NHCONHSO ₂		
20	173	n-Pr	- (CH ₂	2)4-	0	-NHCONHSO2		
	174	n-Pr	- (CH ₂	2)4-	0	-OP3H2	F,H	
	175	n-Pr	- (CH ₂	2)4-	0	-0S03H	F,H	l
	176	n-Pr	- (CH ₂	2)4-	0	-so ₂ NHCO (c		
	177	n-Pr	- (CH ₂	2)4-	0	-so ₂ NHCOCH	2 ^C 6 ^H 5 F, H	
25	178	n-Pr	- (CH ₂	2)4-	0	-so ₂ nhco (n		
	179	n-Pr	- (CH ₂	2)4-	0	-so ₂ nhconh		
	180	n-Pr	- (CH ₂	2)4-	0	-so ₂ nHco(i		
	181	n-Pr	- (CH ₂	2)4-	0	-so ₂ nHCo (n		
	182	n-Pr	- (CH ₂	2)4-	0	-so ₂ nhconh		
30	183	n-Pr	- (CH ₂	2)4-	0	-so ₂ nhconh		
	184	n-Pr	- (CH ₂	4-	0	-so ₂ nhch ₂ c	• -	
	185	n-Pr	- (CH ₂		0	-so ₂ nHCOC ₆	-	
	186	n-Pr	- (CH ₂	4-	0	-so ₂ nHco (n		
	187	n-Pr	- (CH ₂	4-	0	-so ₂ nhco (c	y-C ₃ H ₅) H, n	-Pr

	EX.	R6	R ⁷	R8	R	9 _R 10	R13		R^2, R^3	
	188	n-Pr	- (CH	2)4-		0	-so2инсосн	2C6H5	H, n-P	r
	189	n-Pr	- (CH	2)4-		0	-so2NHCO(n	-Bu)	H, n-P	r
	190	n-Pr	- (CH	2)4-		0	-so ₂ nhconh	(n-Bu)	H, n-P	r
5	191	n-Pr	- (CH	2)4-		0	-so ₂ NHCO(i-	-Bu)	H, n-P	r
	192	n-Pr	- (CH	2)4-		0	-so ₂ NH (n-C	5 ^H 11)	H, n-P	r
	193	n-Pr	- (CH	2)4-		0	-so ₂ NH (i-C	5H ₁₁)	H, n-P	r
	194	n-Pr	- (CH	2)4-		0	-so ₂ nh (cy-	C3H5)	H, n-P	r
	195	n-Pr	- (CH	2)4-		0	-so ₂ NHCH ₂ C	5 ^H 5	H, n-P	r
10	196	n-Pr	- (CH	2)4-		0	-so2NHCOC6	¹ 5	Cl, n-	Pr
	197	n-Pr	- (CH	2)4-		0	-so2NHCO(n-	-C5H ₁₁)	Cl, n-1	Pr
	198	n-Pr	- (CH	2)4-		0	-so ₂ NHCO (c)	/-C3H5)	F,n-P:	r
	199	n-Pr	- (CH ₂	2)4-		0	-so ₂ NHCOCH ₂	2C6H5	F,n-P:	r
	200	n-Pr	- (CH2	2)4-		0	-so ₂ NHCO (n-	-Bu)	Cl,n-	Pr
15	201	n-Pr	- (CH ₂	2)4-	(0	-so2NHCOC2H	¹ 5	F,n-P	r
	202	n-Pr	-(CH ₂	2)4-	(0	-so ₂ NHCOCH ₃	3	Cl,n-	?r
	203	n-Pr	- (CH ₂	2)4-		0	-so ₂ NHCO(i-	C5H11)	F,n-P	
	204	n-Pr	-(CH ₂	2)4-	(0	-сн ₂ со ₂ н		F,n-P	•
	205	n-Pr	- (CH ₂	2)4-	(0	-conhnhso2	F ₃	Cl,n-E	r
20	206	n-Bu	-(CH ₂	2)4-	(>	-coneso2 (n-	C5H11)	Н,Н	
	207	n-Bu	- (CH ₂	2)4-	()	-so ₂ NHCO (cy	-C ₃ н ₅)	н,н	
	208	n-Bu	- (CH ₂	2)4-	()	-so ₂ NHCOCH ₂	C6H5	н,н	
	209	n-Bu	- (CH ₂	2) 4-	()	-coneso2 (n-	·Bu)	н, н	
	210	n-Bu	-(CH ₂	2)4-	()	-conhso ₂ (i-	Bu)	H,H	
25	211	n-Bu	- (CH ₂	4-	()	-conhso ₂ (n-	C ₅ H ₁₁)	н, н	
	212	n-Bu	- (CH ₂) 4-	()	-conhso ₂ (i-	C5H11)	н, н	
	213	n-Bu	-(CH ₂) 4-	c)	-conhso ₂ (cy	-C3H5)	н,н	
	214	n-Bu	- (CH ₂	-	C)	-so ₂ NHCOCH ₂	C6H5	н,н	
	215	n-Bu	-(CH ₂) 4-	C)	-so ₂ NHCOC ₆ H	5	сн3,н	
30	216	n-Bu	-(CH ₂) 4-	C		-so2NHCO(n-	C ₅ H ₁₁)	сн3,н	
	217	n-Bu	- (CH ₂) 4-	C)	-so ₂ nhco(cy	-c ₃ H ₅)	сн3,н	
	218	n-Bu	- (CH ₂		c)	-so ₂ NHCOCH ₂	C6H5	CH3,H	
	219	n-Bu	-(CH ₂) 4-	C)	-NHCONHSO ₂ (n-Bu)	с н 3,н	
	220	n-Bu	-(CH ₂) ₄ -	c)	-NHCONHSO ₂ (i-Bu)	сн ₃ ,н	

	EX. R ⁶	R ⁷ R ⁸	R9R10	R ¹³	\mathbb{R}^2 , \mathbb{R}^3
	221 n-Bu	- (CH ₂) ₄ -	0	-SO2NHCONH(i-Bu)	сн3,н
	222 n-Bu	-(CH ₂) ₄ -	0	$-so_2NH (n-c_5H_{11})$	сн3,н
	223 n-Bu	-(CH ₂) ₄ -	0	$-so_2$ NH (i- c_5 H ₁₁)	сн3,н
5	224 n-Bu	-(CH ₂) ₄ -	o	$-so_2NH_2 (cy-C_3H_5)$	сн3,н
	225 n-Bu	- (CH ₂) ₄ -	o	-so2nHCH2C6H5	сн3,н
	226 n-Bu	-(CH ₂) ₄ -	0	-so ₂ nhcoc ₆ h ₅	Cl,H
	227 n-Bu	-(CH ₂) ₄ -	0	-so ₂ NHCO(n-C ₅ H ₁₁)	Cl,H
	228 n-Bu	-(CH ₂) ₄ -	0	$-so_2$ NHCO(cy- c_3H_5)	Cl,H
10	229 n-Bu	-(CH ₂) ₄ -	0	-so ₂ NHCOCH ₂ C ₆ H ₅	Cl,H
	230 n-Bu	-(CH ₂) ₄ -	0	-NHSO2NHCO(n-Bu)	Cl,H
	231 n-Bu	-(CH ₂) ₄ -	0	-SO2NHCONH (n-Bu)	Cl,H
	232 n-Bu	-(CH ₂) ₄ -	0	-NHSO2NHCO(i-Bu)	Cl,H
	233 n-Bu	-(CH ₂) ₄ -	0	$\verb -NHSO _2 NHCO(n-C_5 H_{11})$	Cl,H
15	234 n-Bu	-(CH ₂) ₄ -	0	$-\text{NHSO}_2\text{NHCO}\left(i-c_5H_{11}\right)$	Cl,H
	235 n-Bu	-(CH ₂) ₄ -	0	-CH ₂ (5-Tetrazoyl)	Cl,H
	236 n-Bu	-(CH ₂) ₄ -	0	-CONH(5-Tetrazoyl)	Cl,H
	237 n-Bu	- (CH ₂) ₄ -	0	-so ₂ NHCOC ₆ H ₅	F,H
	238 n-Bu	-(CH ₂) ₄ -	0	-so ₂ NHCO (n-C ₅ H ₁₁)	F,H
20	239 n-Bu	-(CH ₂) ₄ -	0	$-so_2$ NHCO (cy $-c_3H_5$)	F,H
	240 n-Bu	-(CH ₂) ₄ -	0	$-so_2$ NHCOCH $_2$ C $_6$ H $_5$	F,H
	241 n-Bu	-(CH ₂) ₄ -	0	-NHSO2NHCO(n-Bu)	F,H
	242 n-Bu	-(CH ₂) ₄ -	0	-SO2NHCONH (n-Bu)	F,H
	243 n-Bu	-(CH ₂) ₄ -	0	-SO2NHCO(i-Bu)	F,H
25	244 n-Bu	-(CH ₂) ₄ -	0	$-so_2NH(n-c_5H_{11})$	F,H
	245 n-Bu	-(CH ₂) ₄ -	0	$-so_2$ NH(i- c_5 H ₁₁)	F,H
	246 n-Bu	-(CH ₂) ₄ -	0	$-so_2$ NH(cy-C ₃ H ₅)	F,H
	247 n-Bu	-(CH ₂) ₄ -	0	-so ₂ nhch ₂ c ₆ h ₅	F,H
	248 n-Bu	-(CH ₂) ₄ -	0	-so ₂ NHCOC ₆ H ₅	H,n-Pr
30	249 n-Bu	-(CH ₂) ₄ -	0	-so ₂ NHCO(n-C ₅ H ₁₁)	H, n-Pr
	250 n-Bu	-(CH ₂) ₄ -	0	-so ₂ nhco (cy-c ₃ H ₅)	H, n-Pr
	251 n-Bu	-(CH ₂) ₄ -	0	-so ₂ NHCOCH ₂ C ₆ H ₅	H, n-Pr
	252 n-Bu	-(CH ₂) ₄ -	0	-CONHSO2NHC2H4Cl	H,n-Pr
	253 n-Bu	- (CH ₂) 4-	0	-CONHSO ₂ (i-Bu)	H,n-Pr

	EX. R ⁶	R ⁷ R ⁸	R9R10	R ¹³	R^2, R^3
	254 n-Bu	-(CH ₂) ₄ -	0	$-conhso_2(n-c_5H_{11})$	H,n-Pr
	255 n-Bu	-(CH ₂) ₄ -	0	$-conhso_2(i-c_5H_{11})$	H,n-Pr
	256 n-Bu	-(CH ₂) ₄ -	0	-conhso ₂ (cy-c ₃ H ₅)	H,n-Pr
5	257 n-Bu	-(CH ₂) ₄ -	0	-conhso2ch2c6h2	H,n-Pr
	258 n-Bu	-(CH ₂) ₄ -	0	-so2NHCOC6H5	Cl,n-Pr
	259 n-Bu	-(CH ₂) ₄ -	0	-so ₂ NHCO (n-C ₅ H ₁₁) ·	Cl,n-Pr
	260 n-Bu	-(CH ₂) ₄ -	0	-so2NHCO(cy-C3H5)	F,n-Pr
	261 n-Bu	-(CH ₂) ₄ -	0	-so2NHCOCH2C6H5	F,n-Pr
10	262 n-Bu	-(CH ₂) ₄ -	0	-NHSO2NHCO (n-Bu)	Cl,n-Pr
	263 n-Bu	-(CH ₂) ₄ -	0	-NHSO2NHCO(i-Bu)	F,n-Pr
	264 n-Bu	-(CH ₂) ₄ -	0	-NHSO2NHCO(n-C5H11)	Cl,n-Pr
	265 n-Bu	- (CH ₂) ₄ -	0	$-\text{NHSO}_2\text{NHCO}(i-c_5\text{H}_{11})$	F,n-Pr
	266 n-Bu	-(CH ₂) ₄ -	0	$-so_2$ NHCONH($i-c_5H_{11}$)	F,n-Pr
15	267 n-Bu	-(CH ₂) ₄ -	0	-NHSO2NHCO (cy-C3H5)	F,n-Pr
	268 n-Bu	-(CH ₂) ₄ -	0	-NHSO2NHCOCH2C6H5	Cl,n-Pr
	269 n-Pr	-(CH ₂) ₅ -	0	-conhso2ch2c6H5	н, н
	270 n-Pr	-(CH ₂) ₅ -	0	-so2NHCOC6H5	н,н
	271 n-Pr	-(CH ₂) ₅₋	0	-so ₂ NHCO(n-C ₅ H ₁₁)	н,н
20	272 n-Pr	-(CH ₂) ₅ -	0	$-so_2$ NHCO(cy- c_3H_5)	н,н
	273 n-Pr	-(CH ₂) ₅ -	0	-so2nHCOCH2C6H5	н, н
	280 n-Pr	- (CH ₂) ₅ -	0	-so2NHCOC6H5	Сн ₃ ,н
	281 n-Pr	-(CH ₂) ₅ -	0	-so2NHCO(n-C5H11)	сн ₃ ,н
	282 n-Pr	-(CH ₂) ₅ -	0	$-so_2$ NHCO(cy- c_3H_5)	сн3,н
25	283 n-Pr	-(CH ₂) ₅ -	0	-so2NHCOCH2C6H5	СН3,Н
	284 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCO(n-Bu)	сн3,н
	285 n-Pr	-(CH ₂) ₅ -	0	-SO2NHCONH (n-Bu)	CH3,H
	286 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCO(i-Bu)	СН3,Н
	287 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCO(n-C5H11)	Сн ₃ , н
30	288 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCO(i-C5H11)	CH3,H
	289 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCO(cy-C3H5)	сн3,н
	290 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCOCH2C6H5	Сн3, н
	291 n-Pr	-(CH ₂) ₅ -	0	-so ₂ NHCOC ₆ H ₅	Cl,H
	292 n-Pr	-(CH ₂) ₅ -	0	-so2NHCO(n-C5H11)	Cl,H

	EX. R ⁶	_R 7 _R 8	R9R10	R ¹³	\mathbb{R}^2 , \mathbb{R}^3
	293 n-Pr	- (CH ₂) ₅ -	О	-so2NHCO(cy-C3H5)	Cl,H
	294 n-Pr	-(CH ₂) ₅ -	0	-so ₂ nhcoch ₂ c ₆ h ₅	Cl,H
	295 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCO(n-Bu)	C1,H
5	296 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCO (n-Bu)	Cl,H
	297 n-Pr	-(CH ₂) ₅ -	0	-SO2NHCONH(i-Bu)	Cl,H
	298 n-Pr	-(CH ₂) ₅ -	0	-so ₂ NHCO (n-C ₅ H ₁₁)	Cl,H
	299 n-Pr	-(CH ₂) ₅ -	0	$-\text{NHSO}_2\text{NHCO}\left(i-\text{C}_5\text{H}_{11}\right)$	C1,H
	300 n-Pr	-(CH ₂) ₅ -	0	$-so_2$ NH (cy- c_3 H ₅)	Cl,H
10	301 n-Pr	-(CH ₂) ₅ -	0	-so2NHCH2C6H5	Cl,H
	302 n-Pr	- (CH ₂) ₅ -	0	-so ₂ NHCOC ₆ H ₅	F,H
	303 n-Pr	-(CH ₂) ₅ -	0	-so ₂ NHCO (n-C ₅ H ₁₁)	F,H
	304 n-Pr	- (CH ₂) ₅ -	0	$-so_2$ NHCO(cy- c_3H_5)	F,H
	305 n-Pr	-(CH ₂) ₅ -	0	-so2nHcocH2C6H5	F,H
15	306 n-Pr	-(CH ₂) ₅ -	0	-so ₂ NHCO (n-Bu)	F,H
	307 n-Pr	-(CH ₂) ₅ -	0	-so ₂ NHCONH (n-Bu)	F,H
	308 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCO(i-Bu)	F,H
	309 n-Pr	-(CH ₂) ₅ -	0	$-\mathtt{NHSO}_2\mathtt{NHCO}(\mathtt{n-C}_5\mathtt{H}_{11})$	F,H
	310 n-Pr	-(CH ₂) ₅ -	0	$\verb -NHSO _2 NHCO (i-C_5 H_{11})$	F,H
20	311 n-Pr	-(CH ₂) ₅ -	0	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-C}_3\text{H}_5\right)$	F,H
	312 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCOCH2C6H5	F,H
	313 n-Pr	-(CH ₂) ₅ -	0	-so2NHCOC6H5	H,n-Pr
	314 n-Pr	-(CH ₂) ₅ -	0	-so ₂ NHCO(n-C ₅ H ₁₁)	H,n-Pr
	315 n-Pr	-(CH ₂) ₅ -	0	$-so_2$ NHCO (cy $-c_3H_5$)	H,n-Pr
25	316 n-Pr	-(CH ₂) ₅ -	0	-so ₂ nHcocH ₂ C ₆ H ₅	H,n-Pr
	317 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCO(n-Bu)	H,n-Pr
	318 n-Pr	- (CH ₂) ₅ -	0	-SO2NHCONH (n-Bu)	H,n-Pr
	319 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCO(i-Bu)	H,n-Pr
	320 n-Pr	- (CH ₂) 5-	0	$-NHSO_2NHCO(n-C_5H_{11})$	H,n-Pr
30	321 n-Pr	-(CH ₂) ₅ -	0	$-\text{NHSO}_2\text{NHCO}(i-\text{C}_5\text{H}_{11})$	H,n-Pr
	322 n-Pr	- (CH ₂) ₅ -	0	$-\text{nHSO}_2\text{nHCO}\left(\text{cy-C}_3\text{H}_5\right)$	H,n-Pr
	323 n-Pr	- (CH ₂) ₅ -	0	-NHSO2NHCOCH2C6H5	H,n-Pr
	324 n-Pr	-(CH ₂) ₅ -	0	-so ₂ NHCOC ₆ H ₅	Cl,n-Pr
	325 n-Pr	-(CH ₂) ₅ -	0	-so ₂ NHCO (n-C ₅ H ₁₁)	C1,n-Pr

	EX. R ⁶	_R 7 _R 8	R9R10	R ¹³	R^2, R^3
	326 n-Pr	-(CH ₂) ₅ -	0	$-so_2$ NHCO (cy- c_3 H ₅)	F,n-Pr
	327 n-Pr	-(CH ₂) ₅ -	0	-so ₂ NHCOCH ₂ C ₆ H ₅	F,n-Pr
	328 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCO(n-Bu)	Cl,n-Pr
5	329 n-Pr	-(CH ₂) ₅ -	0	-SO2NHCONH (n-Bu)	Cl,n-Pr
	330 n-Pr	- (CH ₂) ₅ -	0	-NHSO2NHCO(i-Bu)	F,n-Pr
	331 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCO(n-C5H11)	Cl,n-Pr
	332 n-Pr	-(CH ₂) ₅ -	0	-NHSO2NHCO(i-C5H11)	F,n-Pr
	333 n-Pr	- (CH ₂) 5-	0	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-C}_3\text{H}_5\right)$	F,n-Pr
10	334 n-Pr	- (CH ₂) 5-	0	-NHSO2NHCOCH2C6H5	Cl,n-Pr
	335 n-Bu	- (CH ₂) ₅ -	0	-so2NHCOC6H5	н,н
	336 n-Bu	-(CH ₂) ₅ -	0	$-so_2$ NHCO (n-C ₅ H ₁₁)	н, н
	337 n-Bu	- (CH ₂) ₅ -	0	-so ₂ nhco(cy-c ₃ h ₅)	н,н
	338 n-Bu	-(CH ₂) ₅ -	0	-so ₂ nhcoch ₂ c ₆ h ₅	н, н
15	339 n-Bu	-(CH ₂) ₅ -	0	-NHSO2NHCO (n-Bu)	н, н
	340 n-Bu	-(CH ₂) ₅ -	0	-NHSO2NHCO(i-Bu)	н, н
	341 n-Bu	-(CH ₂) ₅ -	0	$-NHSO_2NHCO(n-C_5H_{11})$	н, н
	342 n-Bu	-(CH ₂) ₅ -	0	$-\text{NHSO}_2\text{NHCO}(i-c_5H_{11})$	н, н
	343 n-Bu	-(CH ₂) ₅ -	0	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-C}_3\text{H}_5\right)$	н, н
20	344 n-Bu	-(CH ₂) ₅ -	0	-NHSO2NHCOCH2C6H5	н, н
	345 n-Bu	-(CH ₂) ₅ -	0	-so ₂ NHCOC ₆ H ₅	сн3,н
	346 n-Bu	-(CH ₂) ₅ -	0	$-so_2$ NHCO (n-C ₅ H ₁₁)	СH3, Н
	347 n-Bu	-(CH ₂) ₅ -	0	-so ₂ NHCO(cy-C ₃ H ₅)	сн3,н
	348 n-Bu	-(CH ₂) ₅ -	0	-so ₂ nhcoch ₂ c ₆ h ₅	сн3,н
25	349 n-Bu	-(CH ₂) ₅ -	0	-NHSO2NHCO (n-Bu)	сн3,н
	350 n-Bu	-(CH ₂) ₅ -	0	-SO2NHCONH (n-Bu)	сн3,н
	351 n-Bu	-(CH ₂) ₅ -	0	-NHSO2NHCO(i-Bu)	Сн ₃ ,н
	352 n-Bu	-(CH ₂) ₅ -	0	-NHSO2NHCO(n-C5H11)	сн ₃ ,н
	353 n-Bu	- (сн ₂) ₅ -	0	$-\text{NHSO}_2\text{NHCO}(i-\text{C}_5\text{H}_{11})$	CH3,H
30	354 n-Bu	- (CH ₂) ₅ -	0	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-c}_3\text{H}_5\right)$	сн3,н
	355 n-Bu	- (CH ₂) ₅ -	0	-NHSO2NHCOCH2C6H5	CH3, H
	356 n-Bu	-(CH ₂) ₅ -	0	-so ₂ NHCOC ₆ H ₅	Cl,H
	357 n-Bu	-(CH ₂) ₅ -	0	$-so_2$ NHCO($n-c_5H_{11}$)	Cl,H
	358 n-Bu	-(CH ₂) ₅ -	0	-so ₂ nhco(cy-c ₃ h ₅)	Cl,H

	EX.	R6	R ⁷	R ⁸	R9R10	R ¹³	$\mathbb{R}^2, \mathbb{R}^3$
	359	n-Bu	- (CH	2)5-	0	$-so_2$ NHCOCH $_2$ C $_6$ H $_5$	Cl,H
	360	n-Bu	- (CH	2)5-	0	-NHSO2NHCO (n-Bu)	Cl,H
	361	n-Bu	- (CH	2)5-	0	-NHSO2NHCO(i-Bu)	Cl,H
5	362	n-Bu	- (CH	2)5-	0	$-\text{NHSO}_2\text{NHCO} (n-C_5H_{11})$	Cl,H
	363	n-Bu	- (CH	2)5-	0	-so2NHCONH(n-C5H11)	Cl,H
	364	n-Bu	- (CH	2)5-	0	-NHSO ₂ NHCO ($i-C_5H_{11}$)	Cl,H
	365	n-Bu	- (CH	2)5-	0	-NHSO2NHCO(cy-C3H5)	Cl,H
	366	n-Bu	- (CH	2)5-	0	-NHSO2NHCOCH2C6H5	Cl,H
10	367	n-Bu	- (CH	2)5-	0	-so ₂ NHCOC ₆ H ₅	F,H
	368	n-Bu	- (CH	2)5-	0	$-so_2$ NHCO($n-C_5H_{11}$)	F,H
	369	n-Bu	- (CH	2) ₅ -	0	$-so_2$ NHCO(cy- c_3H_5)	F,H
	370	n-Bu	- (CH	2)5-	0	-so2NHCOCH2C6H5	F,H
	371	n-Bu	- (CH	2)5-	0	-NHSO2NHCO(n-Bu)	F,H
15	372	n-Bu	- (CH	2) ₅ -	0	-NHSO2NHCO(i-Bu)	F,H
	373	n-Bu	- (CH	2)5-	0	$-NHSO_2NHCO(n-C_5H_{11})$	F,H
	374	n-Bu	- (CH	2)5-	0	-so ₂ NHCONH(n-C ₅ H ₁₁)	F,H
	375	n-Bu	- (CH	2)5-	0	$-\text{NHSO}_2\text{NHCO}\left(i-\text{C}_5\text{H}_{11}\right)$	F,H
	376	n-Bu	- (CH	2)5-	0	-NHSO2NHCO(cy-C3H5)	F,H
20	377	n-Bu	- (CH	2)5-	0	-NHSO2NHCOCH2C6H5	F,H
	378	n-Bu	- (CH	₂) ₅ -	0	-so ₂ nhcoc ₆ H ₅	H,n-Pr
	379	n-Bu	- (CH	2)5-	0	$-so_2$ NHCO (n-C ₅ H ₁₁)	H,n-Pr
	380	n-Bu	- (CH	2)5-	0	-so2NHCO(cy-C3H5)	H,n-Pr
	381	n-Bu	- (CH	2)5-	0	-so ₂ nhcoch ₂ c ₆ h ₅	H,n-Pr
25	382	n-Bu	- (CH	2)5-	0	-NHSO2NHCO(n-Bu)	H,n-Pr
	383	n-Bu	- (CH	2)5-	0	-NHSO2NHCO(i-Bu)	H,n-Pr
	384	n-Bu	- (CH	2)5-	0	-so2NHCONH(n-C5H11)	H,n-Pr
	385	n-Bu	- (CH	2)5-	0	$-NHSO_2NHCO(n-C_5H_{11})$	H,n-Pr
	386	n-Bu	- (CH	2)5-	0	$-\text{NHSO}_2\text{NHCO}(i-c_5H_{11})$	H,n-Pr
30	387	n-Bu	- (CH	2)5-	0	-NHSO2NHCO(cy-C3H5)	H,n-Pr
	388	n-Bu	- (CH	2)5-	0	-NHSO2NHCOCH2C6H5	H,n-Pr
	389	n-Bu	- (CH	2)5-	0	-so2nHCOC6H5	Cl,n-Pr
	390	n-Bu	- (CH	2)5-	0	-so ₂ NHCO (n-C ₅ H ₁₁)	Cl,n-Pr
	391	n-Bu	- (CH	2)5-	0	-so2NHCO(cy-C3H5)	F,n-Pr

	EX	. R6	_R 7 _R 8	R9R10	R ¹³	R ² , R ³
	392	n-Bu	-(CH ₂) ₅ -	0	-so ₂ nhcoch ₂ c ₆ h ₅	F,n-Pr
	393	n-Bu	-(CH ₂) ₅ -	0	-NHSO2NHCO(n-Bu)	Cl,n-Pr
	394	n-Bu	-(CH ₂) ₅ -	0	-so ₂ nhconh (n-Bu)	Cl,n-Pr
5	395	n-Bu	-(CH ₂) ₅ -	0	-SO ₂ NHCONH(i-Bu)	Cl,n-Pr
	396	n-Bu	-(CH ₂) ₅ -	0	-NHSO2NHCO(i-Bu)	F,n-Pr
	397	n-Bu	-(CH ₂) ₅ -	0	-NHSO2NHCO (n-C5H11)	Cl,n-Pr
	398	n-Bu	-(CH ₂) ₅ -	0	$-\text{NHSO}_2\text{NHCO}\left(i-\text{C}_5\text{H}_{11}\right)$	F,n-Pr
	399	n-Bu	-(CH ₂) ₅ -	0	-NHSO2NHCO(cy-C3H5)	F, n-Pr
10	400	n-Bu	-(CH ₂) ₅ -	0	-NHSO2NHCOCH2C6H5	Cl,n-Pr
	401	n-Pr	- (CH ₂) S (CH ₂) 2	- 0	-сомнsо ₂ сн ₂ с ₆ н ₅	н, н
	402	n-Pr	-(CH2)S(CH2)2	- 0	-so2nHCOC6H5	н, н
	403	n-Pr	-(CH2)S(CH2)2	- 0	-so ₂ NHCO(n-C ₅ H ₁₁)	Н,Н
	404	n-Pr	-(CH ₂)S(CH ₂) ₂ -	- 0	-so2nhco(cy-c3H5)	н, н
15	405	n-Pr	-(CH2)S(CH2)2	- 0	-so2NHCOCH2C6H5	н, н
	406	n-Pr	-(CH2)S(CH2)2-	- 0	-NHSO2NHCO(n-Bu)	Н,Н
	407	n-Pr	-(CH2)S(CH2)2-	- 0	-NHSO2NHCO(i-Bu)	н, н
	408	n-Pr	- (CH ₂) S (CH ₂) 2	- 0	-NHSO2NHCO (n-C5H11)	н,н
	409	n-Pr	- (CH ₂) S (CH ₂) ₂ -	· o	-NHSO2NHCO(i-C5H11)	н, н
20	410	n-Pr	- (CH ₂) S (CH ₂) 2-	. 0	-NHSO2NHCO (cy-C3H5)	н,н
	411	n-Pr	-(CH2)S(CH2)2-	. 0	-NHSO2NHCOCH2C6H5	н,н
	412	n-Pr	-(CH2)S(CH2)2-	. 0	-so2NHCOC6H5	сн ₃ ,н
	413	n-Pr	- (CH ₂) S (CH ₂) 2-	. 0	$-so_2$ NHCO (n- c_5H_{11})	сн3,н
	414	n-Pr	- (CH ₂) S (CH ₂) 2-	0	-so2NHCO(cy-C3H5)	CH3,H
25	415	n-Pr	-(CH ₂)S(CH ₂) ₂ -	0	-so ₂ nhcoch ₂ c ₆ h ₅	СН3,Н
	416	n-Pr	-(CH ₂)S(CH ₂) ₂ -	0	-NHSO2NHCO(n-Bu)	СН3,Н
	417	n-Pr	- (CH ₂) S (CH ₂) 2-	0	-SO2NHCONH (n-Bu)	сн3,н
	418	n-Pr	- (CH ₂) S (CH ₂) 2-	0	-NHSO2NHCO(i-Bu)	сн3,н
	419	n-Pr	- (CH ₂) S (CH ₂) 2-	0	-NHSO2NHCO (n-C5H11)	CH3,H
30	420	n-Pr	-(CH2)S(CH2)2-	0	$-\text{NHSO}_2\text{NHCO}(i-c_5\text{H}_{11})$	CH3,H
	421	n-Pr	- (CH ₂) s (CH ₂) 2-	0	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-C}_3\text{H}_5\right)$	Сн₃,н
	422	n-Pr	- (CH ₂) s (CH ₂) 2-	0	-NHSO2NHCOCH2C6H5	сн3,н
	423	n-Pr	-(CH2)S(CH2)2-	0	-so2nHcoc6H5	Cl,H
	424	n-Pr	- (CH2) S (CH2) 2-	0	-SO2NHCO(n-C5H11)	Cl,H

	EX. R ⁶	_R 7 _R 8	R9R10	R ¹³	$\mathbb{R}^2, \mathbb{R}^3$
	425 n-Pr	-(CH2)S(CH2)2-	0	$-so_2$ NHCO(cy-C ₃ H ₅)	Cl,H
	426 n-Pr	-(CH2)S(CH2)2-	0	-so ₂ nhcoch ₂ c ₆ H ₅	Cl,H
	427 n-Pr	-(CH2)S(CH2)2-	0	-NHSO2NHCO (n-Bu)	Cl,H
5	428 n-Pr	-(CH2)S(CH2)2-	0	-NHSO2NHCO(n-Bu)	Cl,H
	429 n-Pr	-(CH2)S(CH2)2-	0	-so ₂ NHCONH(i-Bu)	Cl,H
	430 n-Pr	-(CH2)S(CH2)2-	0	$-\text{NHSO}_2\text{NHCO}(n-C_5H_{11})$	C1,H
	431 n-Pr	-(CH2)S(CH2)2-	0	$-\text{NHSO}_2\text{NHCO}(i-c_5H_{11})$	Cl,H
	432 n-Pr	-(CH2)S(CH2)2-	0	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-C}_3\text{H}_5\right)$	Cl,H
10	433 n-Pr	- (CH ₂) S (CH ₂) 2-	0	-NHSO2NHCOCH2C6H5	Cl,H
	434 n-Pr	-(CH2)S(CH2)2-	0	-so2NHCOC6H5	F,H
	435 n-Pr	-(CH ₂) S (CH ₂) ₂ -	0	$-so_2$ NHCO(n-C ₅ H ₁₁)	F,H
	436 n-Pr	-(CH2)S(CH2)2-	0	$-so_2$ NHCO(cy-C ₃ H ₅)	F,H
	437 n-Pr	-(CH2)S(CH2)2-	0	-so2NHCOCH2C6H5	F,H
15	438 n-Pr	-(CH2)S(CH2)2-	0	-NHSO2NHCO(n-Bu)	F,H
	439 n-Pr	-(CH2)S(CH2)2-	0	-SO2NHCONH (n-Bu)	F,H
	440 n-Pr	-(CH2)S(CH2)2-	0	-NHSO2NHCO(i-Bu)	F,H
	441 n-Pr	-(CH2)S(CH2)2-	0	$-\text{NHSO}_2\text{NHCO}(n-C_5\text{H}_{11})$	F,H
	442 n-Pr	-(CH2)S(CH2)2-	0	$-\text{NHSO}_2\text{NHCO}(i-\text{C}_5\text{H}_{11})$	F,H
20	443 n-Pr	-(CH ₂)S(CH ₂) ₂ -	0	-NHSO2NHCO(cy-C3H5)	F,H
	444 n-Pr	-(CH2)S(CH2)2-	0	-NHSO2NHCOCH2C6H5	F,H
	445 n-Pr	-(CH2)S(CH2)2-	0	-so2NHCOC6H5	H,n-Pr
	446 n-Pr	$-(CH_2) S (CH_2)_2-$	0	-so ₂ NHCO(n-C ₅ H ₁₁)	H,n-Pr
	447 n-Pr	-(CH2)S(CH2)2-	0	$-so_2$ NHCO(cy-C ₃ H ₅)	H,n-Pr
25	448 n-Pr	- (CH ₂) S (CH ₂) 2-	0	-so2NHCOCH2C6H5	H,n-Pr
	449 n-Pr	-(CH2)S(CH2)2-	0	-NHSO2NHCO (n-Bu)	H,n-Pr
	450 n-Pr	-(CH ₂) s(CH ₂) ₂ -	0	-SO2NHCONH(n-Bu)	H,n-Pr
	451 n-Pr	- (CH ₂) S (CH ₂) 2-	0	-NHSO2NHCO(i-Bu)	H,n-Pr
	452 n-Pr	- (CH ₂) S (CH ₂) 2-	0	$-NHSO_2NHCO(n-C_5H_{11})$	H,n-Pr
30	453 n-Pr	- (CH ₂) S (CH ₂) 2-	0	$-\text{NHSO}_2\text{NHCO}(i-C_5H_{11})$	H,n-Pr
	454 n-Pr	- (CH ₂) S (CH ₂) 2-	0	-NHSO2NHCO (cy-C3H5)	H,n-Pr
	455 n-Pr	$-$ (CH ₂) S (CH ₂) $_2$ -	0	-NHSO2NHCOCH2C6H5	H,n-Pr
	456 n-Pr	- (CH ₂) S (CH ₂) 2-	0	-so ₂ NHCOC ₆ H ₅	Cl, n-P
	457 n-Pr	$-(CH_2) S (CH_2)_2-$	0	$-so_2$ NHCO (n-C ₅ H ₁₁)	Cl, n-P

	EX	. R ⁶	R ⁷	R ⁸	R9R10	R ¹³	R^2, R^3
	458	n-Pr	-(CH ₂) S (CH ₂) 2-	0	$-so_2$ NHCO (cy- C_3H_5)	F,n-Pr
	459	n-Pr	- (CH ₂) S (CH ₂) 2-	0	-so2nHCOCH2C6H5	F,n-Pr
	460	n-Pr	- (CH ₂) S (CH ₂) 2-	0	-NHSO2NHCO (n-Bu)	Cl,n-Pr
5	461	n-Pr	- (CH ₂) S (CH ₂) 2-	0	-SO2NHCONH (n-Bu)	Cl,n-Pr
	462	n-Pr	- (CH ₂) S (CH ₂) 2-	0	-NHSO2NHCO(1-Bu)	F,n-Pr
	463	n-Pr	-(CH ₂) S (CH ₂) 2-	0	-NHSO2NHCO(n-C5H11)	Cl,n-Pr
	463	n-Pr	-(CH ₂) S (CH ₂) 2-	0	$-\text{NHSO}_2\text{NHCO}(i-\text{C}_5\text{H}_{11})$	F,n-Pr
	464	n-Pr	-(CH ₂) S (CH ₂) 2-	0	-NHSO2NHCO(cy-C3H5)	F,n-Pr
10	465	n-Pr	-(CH ₂) S (CH ₂) 2-	0	-NHSO2NHCOCH2C6H5	Cl,n-Pr
	466	n-Bu	-(CH ₂) S (CH ₂) 2-	0	-so2nHCOC6H5	Н,Н
	467	n-Bu	-(CH ₂) S (CH ₂) ₂ -	0	-so ₂ NHCO (n-C ₅ H ₁₁)	н, н
	468	n-Bu	-(CH ₂) S (CH ₂) ₂ -	0	-so ₂ NHCO(cy-C ₃ H ₅)	н, н
	469	n-Bu	-(CH ₂) S (CH ₂) 2-	0	-so2NHCOCH2C6H5	н, н
15	470	n-Bu	-(CH ₂) S (CH ₂) 2-	0	-NHSO2NHCO (n-Bu)	н, н
	471	n-Bu	-(CH ₂) S (CH ₂) 2-	0	-NHSO2NHCO(i-Bu)	н, н
	472	n-Bu	-(CH ₂) S (CH ₂) 2-	0	-NHSO2NHCO (n-C5H11)	н, н
	473	n-Bu	-(CH ₂) S (CH ₂) 2-	0	$-nHso_2nHco(i-c_5H_{11})$	н, н
	474	n-Bu	- (CH ₂) S (CH ₂) 2-	0	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-C}_3\text{H}_5\right)$	н, н
20	475	n-Bu	-(CH ₂) S (CH ₂) 2-	0	-NHSO2NHCOCH2C6H5	H,H
	476	n-Bu	- (CH ₂) S (CH ₂) 2-	0	-so2NHCOC6H5	CH3,H
	477	n-Bu	- (CH ₂) S (CH ₂) 2-	0	$-so_2$ NHCO ($n-c_5H_{11}$)	сн3,н
	478	n-Bu	- (CH ₂) S (CH ₂) 2-	0	$-so_2$ NHCO(cy- c_3H_5)	CH3,H
	479	n-Bu	-(CH ₂) S (CH ₂) 2-	0	-so ₂ nhcoch ₂ c ₆ H ₅	сн3,н
25	480	n-Bu	- (CH ₂) S (CH ₂) 2-	0	-NHSO2NHCO (n-Bu)	сн3,н
	481	n-Bu	- (CH ₂)) S (CH ₂) ₂ -	0	-SO2NHCONH (n-Bu)	CH3,H
	482	n-Bu	- (CH ₂)	S (CH ₂) ₂ -	0	-NHSO2NHCO(i-Bu)	сн3,н
	483	n-Bu	-(CH ₂	S (CH ₂) ₂ -	0	$-\text{NHSO}_2\text{NHCO} (n-c_5\text{H}_{11})$	сн3,н
	484	n-Bu	-(CH ₂)	S (CH ₂) ₂ -	0	$-\text{NHSO}_2\text{NHCO}(i-c_5\text{H}_{11})$	CH3,H
30	485	n-Bu	- (CH ₂)	S(CH ₂) ₂ -	0	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-C}_3\text{H}_5\right)$	сн3,н
	486	n-Bu	-(CH ₂)	S (CH ₂) ₂ -	0	$-\mathtt{NHSO_2NHCOCH_2C_6H_5}$	CH3,H
	487	n-Bu	-(CH ₂)	s (CH ₂) 2-	0	-so2NHCOC6H5	Cl,H
	488	n-Bu	-(CH ₂)	S (CH ₂) ₂ -	0	$-so_2$ NHCO(n- C_5 H ₁₁)	Cl,H
	489	n-Bu	-(CH ₂)	S (CH ₂) ₂ -	0	-so2NHCO(cy-C3H5)	Cl,H

	EX. R ⁶	_R 7 _R 8	R9R10	R ¹³	R^2, R^3
	490 n-Bu	-(CH2)S(CH2)2-	• о	-so ₂ nhcoch ₂ c ₆ h ₅	Cl,H
	491 n-Bu	- (CH ₂) S (CH ₂) 2	. 0	-NHSO2NHCO(n-Bu)	Cl,H
	492 n-Bu	- (CH ₂) S (CH ₂) 2-	. 0	-NHSO2NHCO(i-Bu)	Cl,H
5	493 n-Bu	- (CH ₂) S (CH ₂) 2-	. 0	-NHSO2NHCO(n-C5H11)	C1,H
	494 n-Bu	- (CH ₂) S (CH ₂) 2	- о	-so ₂ NHCONH(n-C ₅ H ₁₁)	Cl,H
	495 n-Bu	- (CH ₂) S (CH ₂) 2-	. 0	$-\text{NHSO}_2\text{NHCO}(i-\text{C}_5\text{H}_{11})$	Cl,H
	496 n-Bu	- (CH2) S (CH2) 2-	. 0	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-C}_3\text{H}_5\right)$	Cl,H
	497 n=Bu	-(CH2)S(CH2)2-	• о	-NHSO2NHCOCH2C6H5	Cl,H
10	498 n-Bu	- (CH ₂) S (CH ₂) 2-	• о	-so ₂ NHCOC ₆ H ₅	F,H
	499 n-Bu	- (CH ₂) S (CH ₂) 2-	. 0	-so ₂ NHCO(n-C ₅ H ₁₁)	F,H
	500 n-Bu	- (CH ₂) S (CH ₂) 2-	. 0	$-so_2$ NHCO(cy- C_3H_5)	F,H
	501 n-Bu	- (CH2) S (CH2) 2-	0	-so ₂ NHCOCH ₂ C ₆ H ₅	F,H
	502 n-Bu	- (CH2) S (CH2) 2-	0	-NHSO2NHCO(n-Bu)	F,H
15	503 n-Bu	- (CH ₂) S (CH ₂) 2-	0	-NHSO2NHCO(i-Bu)	F,H
	504 n-Bu	- (CH2) S (CH2) 2-	0	-NHSO2NHCO(n-C5H11)	F,H
	505 n-Bu	- (CH ₂) S (CH ₂) 2-	0	$-so_2$ NHCONH (n-C5H ₁₁)	F,H
	506 n-Bu	- (CH ₂) S (CH ₂) 2-	0	$-\text{NHSO}_2\text{NHCO}\left(i-\text{C}_5\text{H}_{11}\right)$	F,H
	507 n-Bu	- (CH ₂) S (CH ₂) 2-	0	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-C}_3\text{H}_5\right)$	F,H
20	508 n-Bu	- (CH ₂) S (CH ₂) 2-	0	-NHSO2NHCOCH2C6H5	F,H
	509 n-Bu	-(CH2)S(CH2)2-	0	-so2NHCOC6H5	H,n-Pr
	510 n-Bu	-(CH2)S(CH2)2-	0	$-so_2$ NHCO(n-C ₅ H ₁₁)	H, n-Pr
	511 n-Bu	-(CH2)S(CH2)2-	0	-so ₂ nhco (cy-C ₃ H ₅)	H,n-Pr
	512 n-Bu	- (CH ₂) S (CH ₂) 2-	0	-so2NHCOCH2C6H5	H, n-Pr
25	513 n-Bu	- (CH ₂) s (CH ₂) ₂ -	0	-NHSO2NHCO (n-Bu)	H,n-Pr
	514 n-Bu	-(CH2)S(CH2)2-	0	-NHSO2NHCO(i-Bu)	H,n-Pr
	515 n-Bu	-(CH2)S(CH2)2-	0	$-so_2$ NHCONH (n-C ₅ H ₁₁)	H, n-Pr
	516 n-Bu	-(CH2)S(CH2)2-	0	$-NHSO_2NHCO(n-C_5H_{11})$	H,n-Pr
	517 n-Bu	- (CH ₂) S (CH ₂) 2-	0	$-\text{NHSO}_2\text{NHCO}\left(i-\text{C}_5\text{H}_{11}\right)$	H, n-Pr
30	518 n-Bu	-(CH ₂)S(CH ₂)2-	o	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-C}_3\text{H}_5\right)$	H, n-Pr
	519 n-Bu	- (CH ₂) S (CH ₂) 2-	0	-NHSO2NHCOCH2C6H5	H,n-Pr
	520 n-Bu	-(CH2)S(CH2)2-	О	-so ₂ NHCOC ₆ H ₅	Cl,n-Pr
	521 n-Bu	-(CH2)S(CH2)2-	0	-so2NHCO(n-C5H11)	C1,n-Pr
	522 n-Bu	- (CH2) S (CH2) 2-	0	-so ₂ NHCO (cy-C ₃ H ₅)	F,n-Pr

	EX.	R6	R ⁷	R8	R9R10	R ¹³	R^2, R^3
	523	n-Bu	- (CH ₂) S (CH ₂) 2	- o	$-so_2$ NHCOCH $_2$ C $_6$ H $_5$	F,n-Pr
	524	n-Bu	- (CH ₂) S (CH ₂) 2	- 0	-NHSO2NHCO(n-Bu)	Cl,n-Pr
	525	n-Bu	- (CH ₂) s (CH ₂) 2-	- 0	-so ₂ NHCONH (n-Bu)	Cl,n-Pr
5	526	n-Bu	- (CH ₂) S (CH ₂) 2-	- 0	-SO ₂ NHCONH(i-Bu)	Cl,n-Pr
	527	n-Bu	-(CH ₂) S (CH ₂) 2-	- 0	-NHSO2NHCO(i-Bu)	F,n-Pr
	528	n-Bu	- (CH ₂) S (CH ₂) 2	- 0	$-\text{NHSO}_2\text{NHCO}(n-c_5H_{11})$	Cl,n-Pr
	529	n-Bu	-(CH ₂) S (CH ₂) 2-	- 0	$-\text{NHSO}_2\text{NHCO}\left(i-c_5H_{11}\right)$	F,n-Pr
	530	n-Bu	- (CH ₂) S (CH ₂) 2-	- 0	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-C}_3\text{H}_5\right)$	F, n-Pr
10	531	n-Bu	- (CH ₂) S (CH ₂) 2-	- 0	-NHSO2NHCOCH2C6H5	Cl,n-Pr
	532	n-Pr	- (CH ₂) o (CH ₂) ₂ -	. 0	-conhso2ch2c6h5	н, н
	533	n-Pr	- (CH ₂) O (CH ₂) 2-	- 0	-so ₂ nHcoc ₆ H ₅	н, н
	534	n-Pr	-(CH ₂) o (CH ₂) 2-	. 0	-so ₂ NHCO (n-C ₅ H ₁₁)	н, н
	535	n-Pr	-(CH ₂) o (CH ₂) 2-	. 0	-so ₂ NHCO (cy-c ₃ H ₅)	н,н
15	536	n-Pr	-(CH ₂) 0 (CH ₂) 2-	. 0	-so ₂ NHCOCH ₂ C ₆ H ₅	н,н
	537	n-Pr	-(CH ₂) O (CH ₂) 2-	. 0	-NHSO2NHCO (n-Bu)	н, н
	538	n-Pr	-(CH ₂) O (CH ₂) 2-	. 0	-NHSO2NHCO(i-Bu)	н, н
	539	n-Pr	-(CH ₂) 0 (CH ₂) 2-	0	-NHSO2NHCO(n-C5H11)	н, н
	540	n-Pr	- (CH ₂) o (CH ₂) ₂ -	. 0	$-\text{NHSO}_2\text{NHCO}\left(i-c_5H_{11}\right)$	н, н
20	541	n-Pr	-(CH ₂) O (CH ₂) 2-	0	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-C}_3\text{H}_5\right)$	н, н
	542	n-Pr	-(CH ₂) O (CH ₂) 2-	0	-NHSO2NHCOCH2C6H5	н,н
	543	n-Pr	-(CH ₂) O (CH ₂) 2-	0	-conhso2ch2c6h2	н,н
	544	n-Pr	-(CH ₂) o (CH ₂) 2-	0	-so2NHCOC6H5	Н,Н
	545	n-Pr	-(CH ₂	0 (CH ₂) 2-	0	$-so_2$ NHCO($n-c_5H_{11}$)	н, н
25	546	n-Pr	- (CH ₂	0 (CH ₂) ₂ -	0	$-so_2$ NHCO(cy- C_3H_5)	н,н
	547	n-Pr	- (CH ₂)	о (СH ₂) 2-	0	-so ₂ NHCOCH ₂ C ₆ H ₅	н,н
	548	n-Pr	-(CH ₂)	0 (CH ₂) ₂ -	0	-NHSO2NHCO(n-Bu)	н, н
	549	n-Pr	- (CH ₂)	0 (CH ₂) 2-	0	-NHSO2NHCO(i-Bu)	н, н
	550	n-Pr	- (CH ₂)	0 (CH ₂) ₂ -	0	$-NHSO_2NHCO(n-C_5H_{11})$	н, н
30	551	n-Pr	- (CH ₂)	0 (CH ₂) 2-	0	$-\text{NHSO}_2\text{NHCO}(i-c_5\text{H}_{11})$	н, н
	552	n-Pr	-(CH ₂)	o (CH ₂) ₂ -	0	-NHSO2NHCO(cy-C3H5)	н,н
	553	n-Pr	- (CH ₂)	0 (CH ₂) 2-	0	$-$ инсосн $_2$ с $_6$ н $_5$	н, н
	554	n-Pr	- (CH ₂) SC	O2 (CH2) 2-	0	-conhso2ch2c6h2	н,н
	555	n-Pr	-(CH ₂)SC	2 (CH ₂) ₂ -	0	-so ₂ NHCOC ₆ H ₅	н,н

	EX.	R6	R ⁷	R 8	R9R10	R ¹³	\mathbb{R}^2 , \mathbb{R}^3
	556	n-Pr	- (CH ₂)	so ₂ (CH ₂) ₂ -	0	$-so_2$ NHCO(n-C ₅ H ₁₁)	н, н
	557	n-Pr	- (CH ₂)	so ₂ (CH ₂) ₂ -	О	$-so_2$ NHCO(cy- c_3H_5)	н, н
	558	n-Pr	- (CH ₂)	so2 (CH2) 2-	О	-so ₂ NHCOCH ₂ C ₆ H ₅	н, н
5	559	n-Pr	- (CH ₂)	so2 (CH2)2-	0	-NHSO2NHCO(n-Bu)	н, н
	560	n-pr	- (CH ₂)	so ₂ (CH ₂) ₂ -	0	-NHSO2NHCO(i-Bu)	н, н
	561	n-Pr	-(CH ₂)	so ₂ (CH ₂) ₂ -	0	$-\text{NHSO}_2\text{NHCO}\left(\text{n-C}_5\text{H}_{11}\right)$	н, н
	562	n-Pr	-(CH ₂)	so ₂ (CH ₂) ₂ -	0	$-\text{NHSO}_2\text{NHCO}(i-\text{C}_5\text{H}_{11})$	н, н
	563	n-Pr	- (CH ₂)	so ₂ (CH ₂) ₂ -	0	-NHSO2NHCO(cy-C3H5)	н,н
10	564	n-Pr	-(CH ₂)	so ₂ (CH ₂) ₂ -	0	-NHSO2NHCOCH2C6H5	н, н
	565	n-Pr	- (CH ₂) NO	COMe (CH ₂) 2	. 0	-conhso2ch2c6H2	н, н
	566	n-Pr	- (CH ₂) NO	COMe (CH ₂) 2-	. 0	-so2NHCOC6H5	н,н
	567	n-Pr	- (CH ₂) NO	COMe (CH ₂) ₂ -	0	$-so_2$ NHCO (n-C ₅ H ₁₁)	н,н
	568	n-Pr	- (CH ₂) NO	COMe (CH ₂) ₂ -	0	-so ₂ NHCO(cy-C ₃ H ₅)	н, н
15	569	n-Pr	-(CH ₂)NC	COMe (CH ₂) ₂ -	0	-so2NHCOCH2C6H5	н,н
	570	n-Pr	- (CH ₂) NO	COMe (CH ₂) ₂ -	0	-NHSO2NHCO (n-Bu)	н,н
	571	n-Pr	- (CH ₂) NO	OMe (CH ₂) ₂ -	0	-NHSO2NHCO(i-Bu)	н, н
	572	n-Pr	- (CH ₂) NO	COMe (CH ₂) ₂ -	0	$-NHSO_2NHCO(n-C_5H_{11})$	н, н
	573	n-Pr	- (CH ₂) NO	OMe (CH ₂) ₂ -	0	-NHSO2NHCO(1-C5H11)	н,н
20	574	n-Pr	- (CH ₂) NO	OMe (CH ₂) ₂ -	0	-NHSO2NHCO(cy-C3H5)	н, н
	575	n-Pr	- (CH ₂) NC	OMe (CH ₂) ₂ -	0	-NHSO2NHCOCH2C6H5	н, н
	576	n-Pr	- (CH ₂) NC	OPh (CH ₂) ₂ -	0	-conhso2ch2c6h2	н, н
	577	n-Pr	- (CH ₂) NC	OPh (CH ₂) ₂ -	0	-so2NHCOC6H5	н, н
	578	n-Pr	- (CH ₂) NC	OPh (CH ₂) 2-	0	-so ₂ NHCO(n-C ₅ H ₁₁)	н, н
25	579	n-Pr	- (CH ₂) NC	OPh (CH ₂) ₂ -	0	$-so_2$ NHCO(cy- c_3H_5)	н, н
	580	n-Pr	- (CH ₂) NC	OPh (CH ₂) ₂ -	0	-so ₂ NHCOCH ₂ C ₆ H ₅	н, н
	581	n-Pr	- (CH ₂) NC	OPh (CH ₂) ₂ -	0	-NHSO2NHCO(n-Bu)	н, н
	582	n-Pr	- (CH ₂) NC	oph (CH ₂) ₂ -	0	-NHSO2NHCO(i-Bu)	н,н
	583	n-Pr	- (CH ₂) NC	OPh (CH ₂) ₂ -	0	-NHSO2NHCO(n-C5H11)	н,н
30	584	n-Pr	- (CH ₂) NC	OPh (CH ₂) ₂ -	0	-NHSO ₂ NHCO(i-C ₅ H ₁₁)	н,н
	585	n-Pr	-	OPh (CH ₂) ₂ -	0	-NHSO2NHCO(cy-C3H5)	н, н
	586	n-Pr	_	OPh (CH ₂) ₂ -	0	-NHSO2NHCOCH2C6H5	н,н
	587	n-Pr	_	H ₂ Ph (CH ₂) ₂		-conhso ₂ ch ₂ c ₆ h ₅	н, н
	588	n-Pr	- (CH ₂) NC	H ₂ Ph (CH ₂) ₂	- 0	-so ₂ NHCOC ₆ H ₅	н,н

	EX.	. R ⁶	R ⁷	R ⁸	$R^{9}R^{10}$	R ¹³	R^2, R^3
	589	n-Pr	- (CH ₂) NC	H ₂ Ph (CH ₂)	₂ - o	$-so_2$ NHCO(n-C ₅ H ₁₁)	н, н
	590	n-Pr	- (CH ₂) NC	H ₂ Ph (CH ₂)	2- 0	$-so_2$ NHCO(cy- c_3H_5)	н, н
	591	n-Pr	-(CH ₂)NC	H ₂ Ph (CH ₂)	₂ - o	-so ₂ nhcoch ₂ c ₆ h ₅	н,н
5	592	n-Pr	-(CH ₂)NC	H ₂ Ph (CH ₂)	₂ - o	-NHSO2NHCO(n-Bu)	н, н
	593	n-Pr	-(CH ₂)NC	H ₂ Ph (CH ₂)	₂ - o	-NHSO2NHCO(i-Bu)	н,н
	594	n-Pr	-(CH ₂)NC	H ₂ Ph (CH ₂)	2- 0	-NHSO2NHCO (n-C5H11)	н, н
	595	n-Pr	- (CH ₂) NC	H ₂ Ph (CH ₂)	2- 0	$-\text{NHSO}_2\text{NHCO}(i-\text{C}_5\text{H}_{11})$	н,н
	596	n-Pr	- (CH ₂) NC	H ₂ Ph (CH ₂)	2- 0	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-C}_3\text{H}_5\right)$	н,н
10	597	n-Pr	- (CH ₂) NC	H ₂ Ph (CH ₂)	2- 0	-NHSO2NHCOCH2C6H5	н,н
	598	n-Pr	CF ₃	CF ₃	0	-so2NHCOC6H5	н,н
	599	n-Pr	CF ₃	CF ₃	0	-so ₂ NHCO(n-C ₅ H ₁₁)	н,н
	600	n-Pr	CF3	CF ₃	0	-so2nhco(cy-c3H5)	Н, Н
	601	n-Pr	CF3	CF ₃	o	-so2nHcocH2C6H5	н, н
15	602	n-Pr	CF ₃	CF ₃	0	-NHSO2NHCO(n-Bu)	н, н
	603	n-Pr	CF3	CF ₃	0	-NHSO2NHCO(i-Bu)	н, н
	604	n-Pr	CF3	CF ₃	0	$-\text{NHSO}_2\text{NHCO}\left(\text{n-C}_5\text{H}_{11}\right)$	н, н
	605	n-Pr	CF ₃	CF ₃	0	$-\text{NHSO}_2\text{NHCO}\left(i-c_5H_{11}\right)$	н, н
	606	n-Pr	CF ₃	CF ₃	0	-NHSO2NHCO(cy-C3H5)	н, н
20	607	n-Pr	CF ₃	CF ₃	0	-NHSO2NHCOCH2C6H5	н, н
	608	n-Pr	Ph	Ph	0	-so2NHCOC6H5	н, н
	609	n-Pr	Ph	Ph	0	$-so_2$ NHCO (n- c_5H_{11})	н, н
	610	n-Pr	Ph	Ph	0	-so2NHCO(cy-C3H5)	н,н
	611	n-Pr	Ph	Ph	0	-so2nhcoch2c6H5	н, н
25	612	n-Pr	Ph	Ph	0	-NHSO2NHCO(n-Bu)	н,н
	613	n-Pr	Ph	Ph	0	-NHSO2NHCO(i-Bu)	н, н
	614	n-Pr	Ph	Ph	0	$-\text{NHSO}_2\text{NHCO}\left(\text{n-C}_5\text{H}_{11}\right)$	H,H
	615	n-Pr	Ph	Ph	0	$-\text{NHSO}_2\text{NHCO}\left(i-c_5H_{11}\right)$	н, н
	616	n-Pr	Ph	Ph	0	$-\text{NHSO}_2\text{NHCO}\left(\text{cy-C}_3\text{H}_5\right)$	н, н
30	617	n-Pr	Ph	Ph	0	$-\text{NHSO}_2\text{NHCOCH}_2\text{C}_6\text{H}_5$	н, н
	618	n-Pr	-(CH ₂) ₂ -	0	-so ₂ NHCOC ₆ H ₅	н,н
	619	n-Pr	- (CH ₂) ₂ -	0	$-so_2$ NHCO (n-C ₅ H ₁₁)	н, н
	620	n-Pr	-(CH ₂) ₂ -	0	-so2nHCO(cy-C3H5)	н,н
	621	n-Pr	-(CH ₂) ₂ -	0	-so2nhcoch2c6H5	H,H

	EX.	R 6	R ⁷	R8	R9R10	R ¹³	R ² , R
	622	n-Pr	- (CH	2)2-	0	-NHSO2NHCO(n-Bu)	н,н
	623	n-Pr	- (CH	2)2-	0	-NHSO2NHCO(i-Bu)	н, н
	624	n-Pr	- (CH ₂	2)2-	0	-NHSO2NHCO(n-C5H11)	н,н
5	625	n-Pr	- (CH	2)2-	0	-NHSO2NHCO(i-C5H11)	н,н
	626 I	n-Pr	- (CH)2-	0	-NHSO2NHCO(cy-C3H5)	Н,Н
	627 :	n-Pr	- (CH	2)2-	0	-NHSO2NHCOCH2C6H5	H,H
	628 1	n-Pr	CH ₃	CH3	S	-so ₂ NHCOC ₆ H ₅	Н,Н
	629 r	n-Pr	CH ₃	CH ₃	s	-CONHSO2CH2C6H5	н,н
10	630 r	n-Pr	CH ₃	CH ₃	s	-so2NHCO(n-C5H11)	H,H

Utility

Angiotensin II (AII) produces numerous biological responses (e.g., vasoconstriction) through stimulation of its receptors on cell membranes. For the purpose of 15 identifying compounds such as AII antagonists which are capable of interacting with the AII receptor, a ligandreceptor binding assay was utilized for the initial screen. The assay was carried out according to the method described by Chiu, et al., Receptor, 1 33, 20 (1990). In brief, aliquots of a freshly prepared particulate fraction of rat adrenal cortex were incubated with 0.05 nM $[^{125}I]$ AII and varying concentrations of potential AII antagonists in a Tris buffer. After a 1 h incubation the reaction was 25 terminated by addition of cold assay buffer. The bound and free radioactivity were rapidly separated through glass-fiber filters, and the trapped radioactivity was quantitated by scintillation counting. The inhibitory concentration (IC_{50}) of potential AII antagonists which 30 gives 50% displacement of the total specifically bound $[125_{\mbox{\scriptsize I}}]$ AII is presented as a measure of the affinity of such compound for the AII receptor.

Using the assay method described above, the compounds of this invention are found to exhibit an

activity of at least IC50 <10 micromolar, thereby demonstrating and confirming the activity of these compounds as effective AII antagonists.

The potential antihypertensive effects of the compounds of this invention may be demonstrated by administering the compounds to awake rats made hypertensive by ligation of the left renal artery [Cangiano et al., J. Pharmacol, Exp. Ther., 1979, 208, 310]. This procedure increases blood pressure by increasing renin production with consequent elevation of AII levels. Compounds are administered intravenously via a cannula in the jugular vein at 10 mg/kg. Arterial blood pressure is continuously measured directly through a carotid artery cannula and recorded using a pressure transducer and a polygraph. Blood pressure levels after 1.5 treatment are compared to pretreatment levels to determine the antihypertensive effects of the compounds. Using the in vivo methodology described above, the

compounds of this invention are found to exhibit an activity (intravenous) which is 10 mg/kg or less, and/or an activity (oral) which is 100 mg/kg or less, thereby demonstrating and confirming the utility of these compounds as effective agents in lowering blood pressure.

20

25 The compounds of the invention are useful in treating hypertension. They are also of value in the management of acute and chronic congestive heart failure and angina. These compounds may also be expected to be useful in the treatment of primary and secondary 30 hyperaldosteronism; renal diseases such as diabetic nephropathy, glomerulonephritis, glomerular sclerosis, nephrotic syndrome, hypertensive nephrosclerosis, end

stage renal disease, used in renal transplant therapy, and to treat renovascular hypertension, scleroderma,

15

20

25

left ventricular dysfunction, systolic and diastolic dysfunction, diabetic retinopathy and in the management of vascular disorders such as migraine, Raynaud's disease, and as prophylaxis to minimize the atherosclerotic process and neointimal hyperplasia following angioplasty or vascular injury and to retard the onset of type II diabetes. The application of the compounds of this invention for these and similar disorders will be apparent to those skilled in the art.

The compounds of this invention are also useful to treat elevated intraocular pressure and to enhance retinal blood flow and can be administered to patients in need of such treatment with typical pharmaceutical formulations such as tablets, capsules, injectables and the like as well as topical ocular formulations in the form of solutions, ointments, inserts, gels and the like. Pharmaceutical formulations prepared to treat intraocular pressure would typically contain about 0.1% to 15% by weight, preferably 0.5% to 2% by weight, of a compound of this invention. For this use, the compounds of this invention may also be used in combination with other medications for the treatment of glaucoma including choline esterase inhibitors such as physostigmine salicylate or demecarium bromide, parasympathominetic agents such as pilocarpine nitrate, β -adrenergic antagonists such as timolol maleate,

In the management of hypertension and the clinical conditions noted above, the compounds of this invention may be utilized with a pharmaceutical carrier in compositions such as tablets, capsules or elixirs for oral administration, suppositories for rectal administration, sterile solutions or suspensions for

adrenergic agonists such as epinephrine and carbonic

anhydrase inhibitors such as MK-507.

parenteral or intramuscular administration, and the like. The compounds of this invention can be administered to patients (animals and human) in need of such treatment in dosages that will provide optimal 5 pharmaceutical efficacy. Although the dose will vary from patient to patient depending upon the nature and severity of disease, the patient's weight, special diet that is being followed by a patient, concurrent medication, and other factors which those skilled in the 10 art will recognize, the dosage range will generally be about 1 to 1000 mg per patient per day which can be administered in single or multiple doses. Preferably, the dosage range will be about 5 to 500 mg per patient per day; more preferably about 5 to 300 mg per patient 15 per day.

The compounds of this invention can also be administered in combination with other antihypertensives and/or diuretics. For example, the compounds of this invention can be given in combination with diuretics 20 such as hydrochlorothiazide, chlorothiazide, chlorthalidone, methylclothiazide, furosemide, ethacrynic acid, triamterene, amiloride spironolactone and atriopeptin; calcium channel blockers, such as diltiazem, felodipine, nifedipine, amlodipine, 25 nimodipine, isradipine, nitrendipine and verapamil; β adrenergic antagonists such as timolol, atenolol, metoprolol, propanolol, nadolol and pindolol; angiotensin converting enzyme inhibitors such as enalapril, lisinopril, captopril, ramipril, quinapril and zofenopril; remin inhibitors such as A-69729, FK 906 30 and FK 744; α-adrenergic antagonists such as prazosin, doxazosin, and terazosin; sympatholytic agents such as methyldopa, clonidine and quanabenz; atriopeptidase inhibitors (alone or with ANP) such as UK-79300;

serotonin antagonists such as ketanserin; \$\(\lambda_2\)-adrenosine receptor agonists such as CGS 22492C; potassium channel agonists such as pinacidil and cromakalim; and various other antihypertensive drugs including reserpine, minoxidil, guanethidine, hydralazinc hydrochloride and sodium nitroprusside as well as combinations of the above-named drugs. Combinations useful in the management of congestive heart failure include, in addition, compounds of this invention with cardiac stimulants such as dobutamine and xamoterol and phosphodiesterase inhibitors including amrinone and milrinone.

Typically, the individual daily dosages for these combinations can range from about one-fifth of the minimally recommended clinical dosages to the maximum 15 recommended levels for the entities when they are given singly. To illustrate these combinations, one of the angiotensin II antagonists of this invention effective clinically in the 5-500 milligrams per day range can be effectively combined at levels at the 1.0-500 milligrams 20 per day range with the following compounds at the indicated per day dose range; hydrochlorothiazide (6-100 mg), chlorothiazide (125-500 mg), ethacrynic acid (5-200 mg), amiloride (5-20 mg), furosemide (5-80 mg), propranolol (10-480 mg), timolol maleate (1-20 mg), 2.5 methyldopa (125-2000 mg), felodipine (1-20 mg), nifedipine (5-120 mg), nitrendipine (5-60 mg), and diltiazem (30-540 mg). In addition, triple drug combinations of hydrochlorothiazide (5-100 mg) plus amiloride (5-20 mg) plus angiotensin II antagonists of 30 this invention (1-500 mg) or hydrochlorothiazide (5-100 mg) plus timolol maleate (5-60 mg) plus an angiotensin II antagonists of this invention (1-500 mg) or

hydrochlorothiazide (5-200 mg) and nifedipine (5-60 mg)

plus an angiotensin II antagonist of this invention (1-500 mg) are effective combinations to control blood pressure in hypertensive patients. Naturally, these dose ranges can be adjusted on a unit basis as necessary 5 to permit divided daily dosage and, as noted above, the dose will vary depending on the nature and severity of the disease, weight of patient, special diets and other factors.

The active ingredient can be administered orally in 10 solid dosage forms, such as capsules, tablets, and powders, or in liquid dosage forms, such as elixirs syrups, and suspensions. It can also be administered parenterally, in sterile liquid dosage forms.

Gelatin capsules contain the active ingredient and 15 powdered carriers, such as lactose, starch, cellulose derivatives, magnesium stearate, stearic acid, and the like. Similar diluents can be used to make compressed tablets. Both tablets and capsules can be manufactured as sustained release products to provide for continuous release of medication over a period of hours. Compressed tablets can be sugar coated or film coated to mask any unpleasant taste and protect the tablet from the atmosphere, or enteric coated for selective disintegration in the gastrointestinal tract.

20

25

Liquid dosage forms for oral administration can contain coloring and flavoring to increase patient acceptance.

In general, water, a suitable oil, saline, aqueous dextrose (glucose), and related sugar solutions and 30 glycols such as propylene glycol or polyethylene glycols are suitable carriers for parenteral solutions. Solutions for parenteral administration preferably contain a water soluble salt of the active ingredient, suitable stabilizing agents, and if necessary, buffer

substances. Antioxidizing agents such as sodium bisulfite, sodium sulfite, or ascorbic acid, either alone or combined, are suitable stabilizing agents. Also used are citric acid and its salts and sodium EDTA. In addition, parenteral solutions can contain preservatives, such as benzalkonium chloride, methyloropylparaben, and chlorobutanol.

Suitable pharmaceutical carriers are described in Remington's Pharmaceutical Sciences, A. Osol, a standard reference text in this field.

Useful pharmaceutical dosage-forms for administration of the compounds of this invention can be illustrated as follows:

15 <u>Capsules</u>

10

20

25

30

A large number of unit capsules are prepared by filling standard two-piece hard gelatin capsules each with 100 milligrams of powdered active ingredient, 150 milligrams of lactose, 50 milligrams of cellulose, and 6 milligrams magnesium stearate.

Soft Gelatin Capsules

A mixture of active ingredient in a digestible oil such as soybean oil, cottonseed oil or olive oil is prepared and injected by means of a positive displacement pump into gelatin to form soft gelatin capsules containing 100 milligrams of the active ingredient. The capsules are washed and dried.

<u>Tablets</u>

A large number of tablets are prepared by conventional procedures so that the dosage unit is 100 milligrams of active ingredient, 0.2 milligrams of colloidal silicon dioxide, 5 milligrams of magnesium

WO 93/04046 PCT/US92/07022

55

stearate, 275 milligrams of microcrystalline cellulose, 11 milligrams of starch and 98.8 milligrams of lactose. Appropriate coatings may be applied to increase palatability or delay absorption.

5

20

Injectable

A parenteral composition suitable for administration by injection is prepared by stirring 1.5% by weight of active ingredient in 10% by volume

10 propylene glycol. The solution is made to volume with water for injection and sterilized.

Suspension

An aqueous suspension is prepared for oral administration so that each 5 milliliters contain 100 milligrams of finely divided active ingredient, 100 milligrams of sodium carboxymethyl cellulose, 5 milligrams of sodium benzoate, 1.0 grams of sorbitol solution, U.S.P., and 0.025 milliliters of vanillin.

The same dosage forms can generally be used when the compounds of this invention are administered stepwise in conjunction with another therapeutic agent. When the drugs are administered in physical combination, the dosage form and administration route should be

25 selected for compatibility with both drugs.

What is claimed is:

A compound of formula (I) 1.

5 wherein:

R1 is other than in the ortho position and is:

R² is

- 10 (a) н,
 - (b) halo (F, Cl, Br, I),
 - (c) C1-C4 alkyl,
 - C1-C4 alkoxy, (d)
 - C1-C4 acyloxy, (e)
- C1-C4 alkylthio, 15 (f)
 - C1-C4 alkylsulfinyl, (g)
 - (h) C1-C4 alkylsulfonyl,
 - hydroxy (C1-C4) alkyl,
 - aryl (C1-C4) alkyl, (j)
 - -CO₂H, (k)

(i)

- (1) -CN,
- -CONHOR12, (m)
- -SO2NHR21, (n)
- -NH2, (0)
- C1-C4 alkylamino, 25 (p)

(q) C1-C4 dialkylamino, -NHSO2R20, (r)

(s) -NO2.

(t) furyl,

5 arvl; (u)

wherein aryl is phenyl optionally substituted with one or two substituents selected from the group consisting of halo, C1-C4 alkyl, C1-C4 alkoxy, -NO2, -CF3, C1-C4 alkylthio, -OH, -NH2, C1-C4 alkylamino, C1-C4 10 dialkylamino, -CN, -CO₂H, -CO₂CH₃, -CO₂CH₂CH₃, -CO₂benzyl;

R3 is

(a) Η,

(b) halo,

15 (c) C1-C4 alkyl,

> (d) C1-C4 alkoxy,

(e) C1-C4 alkoxyalkyl;

 \mathbb{R}^4 is

(a) -CN,

20 (b) -NO2,

> (c) -CO2R11;

R⁵ is

(a) н.

(b) C1-C6 alkyl,

25 (c) C3-C6 cycloalkyl,

> (d) C2-C4 alkenyl,

(e) C2-C4 alkynyl;

R6 is

30

(a) C_1-C_{10} alkyl,

(b) C3-C8 alkenyl,

(c) C3-C8 alkynyl,

(d) C3-C8 cycloalkyl,

C4-C8 cycloalkenyl, (e)

(f) C4-C10 cycloalkylalkyl,

ε

- (q) C₅-C₁₀ cycloalkylalkenyl,
- (h) C₅-C₁₀ cycloalkylalkynyl,
- (i) (CH₂) sZ (CH₂) mR⁵,
- (j) phenyl, optionally substituted with 1-2
- 5 substituents selected from the group of halo, C_1-C_4 alky1, C_1-C_4 alkoxy, $-NO_2$, $-NH_2$, -OH and benzyloxy,
 - (k) benzyl, optionally substituted on the phenyl ring with 1-2 substituents selected from the group of halo, C_1-C_4 alkyl, C_1-C_4 alkoyy and $-NO_2$;

10

- R7, R8, R9, and R10 are independently chosen from
 - (a) H
 - (b) C₁-C₈ alkyl,
 - (c) C₁-C₈ perfluoroalkyl,
- 15 (d) C₃-C₆ cycloalkyl,
 - (e) -NO₂,
 - (f) -CN,
 - (g) -CONR¹⁵R¹⁶,
 - (h) $-CO_2R^{17}$,
- 20 (i) -OR¹⁸,
 - (j) $-(CH_2)_nCONR^{15}R^{16}$ where n is 1-4,
 - (k) $-(CH_2)_nCO_2R^{17}$ where n is 1-4,
 - (1) $(CH_2)_{p}OR^{18}$ where n is 1-4,
 - (m) aryl, wherein aryl is as defined above,
 - (n) -CH₂ aryl, wherein aryl is as defined above,
 - (o) R^7 and R^8 taken together are $-(CH_2)t^-$, or
 - $-(CH_2)_mX(CH_2)_{q}-,$
 - (p) R^9 and R^{10} taken together can be S, or O;

 \mathbb{R}^{11} is

25

- (a) H,
 - (b) C_1-C_4 alkyl,
 - (c) C₁-C₄ cycloalkyl,
 - (d) phenvl,
 - (e) benzyl;

(a) н,

methyl, (b)

(c) benzyl;

 R^{13} is

(a) -CH2CO2H,

(b) -C (CF3) 2OH,

(c) -CONHNHSO2CF3,

(d) -CONHOR12,

-CONHSO2R20, 10 (e)

> (f) -CONHSO2NHR19,

-C (OH) R19PO3H2, (g)

(h) -NHCONHSO2R20,

-SO2NHCOR19,

(i) -NHPO3H2,

15

(j)

(k) -OPO3H2,

(1) -OSO3H,

-PO (OH) R19, (m)

-PO3H2, (n)

20

(0) -SO3H,

-SO2NHR19, (p)

(q) -NHSO2NHCOR19,

-SO2NHCONHR19, (r)

(s)

(t)

25

(u)

 R^{14} is

- (a) Η,
- (b) C1-C4 alkyl,
 - aryl, wherein aryl is defined as above, (c)
 - (d) benzyl,
 - (e) COR11,
 - (f) CONHR11;
- R15 and R16 are independently 10
 - (a) н,

above,

- (b) C1-C6 alkyl,
- aryl, wherein aryl is as defined above, (c)
- aryl (C_1-C_4) alkyl, where aryl is as defined (d)
- 15

or taken together constitute a

- piperidine ring, (e)
- (f) morpholine ring,
- piperazine ring, optionally N-substituted with (g)
- C1-C6 alkyl, phenyl or benzyl; 20

R17 is

- (a) н,
- C1-C6 alkyl, (b)
- (c) phenyl,
- 25 (d) benzvl;

 R^{18} is

- (a)
- Н, (b) C1-C6 alkyl,
- phenyl, (c)
- (d) benzyl; 30

R19 is

(a) н.

R²⁰ is

- (b) C_1 - C_5 alkyl optionally substituted with a substituent selected from the group consisting of aryl, where aryl is as defined above, heteroaryl where heteroaryl is as defined above, -OH, -SH, C_1 - C_4 alkyl, C_1 - C_4 alkoxy, C_1 - C_4 alkylthio, -CF3, halo, -NO2, -CO2H,
- C₁-C₄ alkoxy, C₁-C₄ alkylthio, -CF₃, halo, -NO₂, -CO₂i -CO₂CH₃, -CO₂-benzyl, -NH₂, C₁-C₄ alkylamino, C₁-C₄ dialkylamino, -PO₃H₂,
 - (c) aryl, where aryl is defined as above,
 - (d) -CH2 aryl, where aryl is defined as above,
- 10 (e) heteroaryl,

wherein heteroaryl is an unsubstituted, monosubstituted or disubstituted 5- or 6-membered aromatic ring which can optionally contain from 1 to 3 heteroatoms selected from the group consisting of O, N, and S and wherein the substituents are members selected from the group consisting of -OH, -SH, C₁-C₄ alkyl, C₁-C₄ alkoxy, -CF₃, halo, -NO₂, -CO₂H, -CO₂CH₃, -CO₂-benzyl, -NH₂, C₁-C₄ alkylamino, C₁-C₄ dialkylamino;

- 20 (a) aryl, where aryl is defined as above,
 - (b) C3-C7 cycloalkyl,
 - (c) C1-C4 perfluoroalkvl.
 - (d) C_1-C_4 alkyl optionally substituted with a
- substituent selected from the group consisting of aryl, 25 where aryl is as defined above, heteroaryl where heteroaryl is as defined above, -OH, -SH, C1-C4 alkyl, C1-C4 alkoxy, C1-C4 alkylthio, -CF3, halo, -NO2, -CO2H, -CO2CH3, -CO2-benzyl, -NH2, C1-C4 alkylamino, C1-C4 dialkylamino, -PO3H2,
- 30 (e) heteroaryl, where heteroaryl is as defined above; R²¹ is
 - (a) H,
 - (b) C₁-C₆ alkyl,

WO 93/04046 PCT/US92/07022

62

ċ

```
(c)
              phenyl,
               benzyl,
         (d)
     or taken together constitute a
              piperidine ring,
         (e)
              morpholine ring,
 5
         (f)
         (g) piperazine ring, optionally N-substituted with
         C1-C6 alkyl, phenyl or benzyl;
    X is
10
         (a) S,
         (b) 0,
         (c) SO,
         (d) SO<sub>2</sub>,
         (e) CHR^{14},
         (f) NR14;
15
     Z is
         (a) -0-,
         (b) -S-,
20
         (c) -NR11-;
    m is 1 to 5;
    n is 1 to 4;
    q is 1 to 5;
    t is 2 to 5;
25
    or a pharmaceutically acceptable salt thereof.
```

 $\label{eq:2.2} \textbf{A compound of Claim 1 wherein} \\ \textbf{30} \quad \textbf{R}^{\textbf{1}} \text{ is in the para position and is}$

R1 is in the para position and is

5

10

20

R6 is

- (a) C1-C10 alkyl,
- (b) C_3-C_{10} alkenyl,
- (c) C₃-C₁₀ alkynyl,
- (d) C3-C8 cycloalkyl,

(e) phenyl, optionally substituted with 1-2 substituents selected from the group of halo, C_1-C_4 alkyl, C_1-C_4 alkoy, $-NO_2$, $-NH_2$, -OH and benzyloxy,

- (f) benzyl, optionally substituted on the phenyl 15 ring with one or two substituents selected from the group consisting of halo, C₁-C₄ alkyl, C₁-C₄ alkoxy and -NO₂;
 - R7, R8, R9, R10 are independently
 - (a) H.
 - (b) C1-C4 alkyl,
 - (c) C₁-C₄ perfluoroalkyl,
 - (d) C3-C6 cycloalkyl,
 - (e) phenyl, optionally substituted with one or two substituents selected from the group of halo, C_1-C_3
- 25 alkyl, C1-C3 alkoxy, NO2, CF3, NH2, and OH,
 - (f) R^7 and R^8 taken together are -(CH2)t-, or -(CH2)mX(CH2)q-,
 - (g) R⁹ and R¹⁰ taken together can be S, O;

WO 93/04046 PCT/US92/07022

64

R¹³ is

- (a) -CONHSO2R20,
- (b) -NHCONHSO2R20,
- (c) -NHSO2NHCOR19,
- (d) -PO₃H₂,
 - (e) -SO3H,
- (f) -SO2NHR19,
- (g) -SO2NHCOR19,
- (h) -SO2NHCONHR19,

10 (i)

5

or a pharmaceutically acceptable salt thereof.

A compound of Claim 2 wherein

15

R6 is

- (a) C_1-C_7 alkyl,
 - (b) C₃-C₄ alkenyl,
 - (c) C3-C4 alkynyl;
- 20 (d) phenyl, optionally substituted with 1-2 substituents selected from the group of halo, C1-C4 alkyl, C1-C4 alkoxy, -NO2, -NH2, -OH and benzyloxy;

R13 is

- 25 (a) -CONHSO₂R²⁰,
 - (b) -NHCONHSO2R20,
 - (c) -SO2NHR19,
 - (d) -SO₂NHCONHR¹⁹,
 - (e) $-SO_2NHCOR^{19}$,
- 30 (f) -SO₂NHCONHR¹⁹;

or a pharmaceutically acceptable salt thereof.

4. A compound of Claim 3 wherein

R¹ is

- or a pharmaceutically acceptable salt thereof.
 - A compound of Claim 4 selected from the group consisting of
- N-((4'-(((4-oxo-2-propyl-1,3-diazaspiro((4.4))non-1-en-3-yl)methyl))((1,1'-biphenyl-2-ylsulfonyl))benzamide
 - N-((4'-(((4-oxo-2-butyl-1,3-diazaspiro((4.4))non-1-en-3-yl)methyl))((1,1'-biphenyl-2-ylsulfonyl))-
- 20 benzamide
 - N-((4'-(4,5-dihydro-4,4-dimethyl-5-oxo-2-propyl-1H-imidazol-1-ylmethyl)((1,1'-biphenyl))-2-ylsulfonyl))-benzamide

- N-((4'-(4,5-dihydro-4,4-dimethyl-5-oxo-2-propyl-1H-imidazol-1-ylmethyl)((1,1'-biphenyl))-2-ylsulfonyl))-4-chlorobenzamide

N-((4'-(((4-oxo-2-propyl-1,3-diazaspiro((2.4)))hept-1-en-3-yl)methyl))((1,1'-biphenyl-2-ylsulfonyl))hexanamide

5

N-((4'-(((4-oxo-2-propyl-1,3-diazaspiro((4.4))non-1-en-3-yl)methyl))((1,1'-biphenyl-2-ylsulfonyl))hexanamide

 N-((4'-(((4-oxo-2-propyl-1,3-diazaspiro((4.5)))dec-10

1-en-3-v1)methvl))((1,1'-biphenvl-2-vlsulfonyl))hexanamide

N-((4'-(((4-oxo-2-butyl-1,3-diazaspiro((4.4))non-1en-3-yl)methyl))((1,1'-biphenyl-2-ylsulfonyl))-15 hexanamide

N-((4'-(4.5-dihydro-4.4-dimethyl-5-oxo-2-propyl-1Himidazol-1-vlmethvl)((1,1'-biphenvl))-2-

20 vlsulfonyl))-hexanamide

> N-((4'-(4,5-dihydro-4,4-dicyclopropyl-5-oxo-2propyl-1H-imidazol-1-ylmethyl) ((1,1'-biphenyl))-2vlsulfonyl))-hexanamide

25

N-((4'-(4,5-dihydro-4,4-bistrifluoromethyl-5-oxo-2propyl-1H-imidazol-1-ylmethyl) ((1,1'-biphenyl))-2ylsulfonyl))-hexanamide

N-((4'-(4,5-dihydro-4,4-dimethyl-5-oxo-2-propyl-1H-30 imidazol-1-ylmethyl)-3'-methyl((1,1'-biphenyl))-2vlsulfonyl))-hexanamide

- N-((4'-(4,5-dihydro-4,4-dimethyl-5-oxo-2-propyl-1H-imidazol-1-ylmethyl)-4-propyl((1,1'-biphenyl))-2-ylsulfonyl))-hexanamide
- 5 N-((4'-(4,5-dihydro-4,4-dimethyl-5-oxo-2-propyl-1H-imidazol-1-ylmethyl)((1,1'-biphenyl))-2-ylsulfonyl))-trifluoroacetamide
- 3,5-dihydro-5,5-dimethyl-2-propyl-3-[(2'-(N-((phenylsulfonyl)carboxamido)biphen-4-yl)methyl]-4Himidazol-4-one.
- A pharmaceutical composition comprising a pharmaceutically suitable carrier and a compound of any 15 one of Claims 1 through 4.
 - A pharmaceutical composition comprising a pharmaceutically suitable carrier and a compound of Claim 5.

8. A method of treating hypertension in a warm blooded animal comprising administering to an animal in need of such treatment an effective amount of a compound of any of Claims 1 through 4.

25

9. A method of treating hypertension in a warm blooded animal comprising administering to an animal in need of such treatment an effective amount of a compound of Claim 5.

30

10. A method of treating congestive heart failure in a warm blooded animal comprising administering to an animal in need of such treatment an effective amount of a compound of any of Claims 1 through 4. WO 93/04046 PCT/US92/07022

68

11. A method of treating congestive heart failure in a warm blooded animal comprising administering to an animal in need of such treatment effective of a compound 5 of Claim 5.

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 92/07022

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) According to International Patent Classification (IPC) nr to both National Classification and IPC A61K31/415: C07D235/02 Int.Cl. 5 C07D233/70; CO7D233/84: C07D495/10 CO7D403/10: CO7D471/10: CO7D491/10: II. FIELDS SEARCHED Minimum Documentation Searched Classification System Classification Symbols C07D ; A61K ; C07F Int.Cl. 5 Documentation Searched other than Minimum Documentation to the Extent that such Documents are included in the Fields Searched III. DOCUMENTS CONSIDERED TO BE RELEVANT⁹ Citation of Document, 11 with indication, where appropriate, of the relevant passages 12 Relevant to Claim Nn.13 Category º | EP,A,O 380 959 (E.I.DU PONT DE NEMOURS AND COMPANY) 8 August 1990 DE, A, 3 913 757 (SCHERING AG) 25 October 1990 EP, A, 0 419 048 (MERCK & CO. INC.) 27 March 1991 cited in the application EP,A,O 412 594 (MERCK & CO. INC.) 13 February 1991 cited in the application EP,A,O 291 969 (E.I.DU PONT DE NEMOURS AND COMPANY) 23 November 1988 cited in the application "T" later document published after the international filling date or priority date and not in conflict with the application but cited in understand the principle or theory underlying the O Special categories of cited documents: 10 "A" document defining the general state of the art which is not considered to be of particular relevance invention "E" earlier document but published nn or after the international "X" document of particular relevance; the claimed loventinn cannot be considered novel or cannot be considered to involve an inventive step filing date "L" document which may throw dnubts no prinrity claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "Y" document in particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combinatina being obvious to a person skilled "O" document referring to an oral discinsure, use, exhibiting or other means "P" document published prior to the international filing date but later than the priority date claimed "&" document member of the same parent family IV. CERTIFICATION Date of Mailing of this International Search Report Date of the Actual Completing of the International Search 1 5, 01, 93 02 DECEMBER 1992 Signature of Authorized Officer international Searching Authority DE BUYSER I.A.F :--EUROPEAN PATENT OFFICE

International Application No

	International Application No	
	INTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)	Relevant to Claim No.
Category °	Citation of Document, with indication, where appropriate, of the relevant passages	
P,X	WO,A,9 114 679 (SANOFI) 3 October 1991 cited in the application see the whole document	1,4-11
E .	cited in the application	1-4,6-11

ANNEX TO THE INTERNATIONAL SEARCH REPORT ON INTERNATIONAL PATENT APPLICATION NO. US SA

9207022 63823

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report. The members are as centained in the European Patent Office EUP file on The European Patent Office is now as paid for the purpose of information. 02/12/92 The European Patent Office is in own justable for these particulars which are merely given for the purpose of information. 02/12/92

Patent document cited in search report	Publication date		Patent family member(s)	Publicatio date
EP-A-0380959	08-08-90	US-A- AU-B- AU-A- CA-A- JP-A-	4916129 618503 4854590 2006604 3002169	10-04-90 19-12-91 26-07-90 19-07-90 08-01-91
DE-A-3913757	25-10-90	None		
EP-A-0419048	27-03-91	US-A- CA-A- JP-A-	5100897 2024113 3197466	31-03-92 01-03-91 28-08-91
EP-A-0412594	13-02-91	CA-A- JP-A-	2021954 3148266	29-01-91 25-06-91
EP-A-0291969	23-11-88	US-A- AU-B- AU-A- JP-A- US-A- US-A-	4820843 603525 1650388 1117876 4870186 4874867	11-04-89 15-11-90 24-11-88 10-05-89 26-09-89 17-10-89
WO-A-9114679	03-10-91	FR-A- FR-A- AU-A- CA-A- EP-A-	2659967 2665702 7561091 2057913 0454511	27-09-91 14-02-92 21-10-91 21-09-91 30-10-91
EP-A-0505098	23-09-92	None		